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I also certify that the attached copy of the request for grant of a Patent (Form 1/77) bears a correction, effected by this office, following a request by the applicant and agreed to by the Comptroller-General.

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Signed

Dated 1 September 2004



26AUG03 E832728-1 003018
P01/7700 0.00-0319917.1

Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

The Patent Office

Cardiff Road
Newport
South Wales
NP10 8QQ

1. Your reference

250/ASW

2. Patent application number

(The Patent Office will fill this part in)

0319917.1

23 AUG 2003

3. Full name, address and postcode of the or of each applicant (underline all surnames)

① BRITISH BIOTECH PHARMACEUTICALS LTD
WATLINGTON ROAD, OXFORD OX4 6LY

② LABORATOIRES SERONO SA
ZONE INDUSTRIELLE L'OURIETTAZ
1170 AUBONNE SWITZERLAND

9373003

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

① ENGLAND AND WALES ② SWITZERLAND

4. Title of the invention

METALLOPROTEINASE INHIBITORS

5. Name of your agent (if you have one)

ADAM J. WALLS

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

BRITISH BIOTECH PHARMACEUTICALS LTD
WATLINGTON ROAD, OXFORD OX4 6LY

9373003.

Patents ADP number (if you know it)

6. Priority: Complete this section if you are declaring priority from one or more earlier patent applications, filed in the last 12 months.

Country

Priority application number

Date of filing

(if you know it)

(day / month / year)

7. Divisionals, etc: Complete this section only if this application is a divisional application or resulted from an entitlement dispute (see note f)

Number of earlier UK application

Date of filing

(day / month / year)

8. Is a Patents Form 7/77 (Statement of inventorship and of right to grant of a patent) required in support of this request?

Answer YES if:

- a) any applicant named in part 3 is not an inventor, or
- b) there is an inventor who is not named as an applicant, or
- c) any named applicant is a corporate body.

Otherwise answer NO (See note d)

YES

Patents Form 1/77

9. Accompanying documents: A patent application must include a description of the invention. Not counting duplicates, please enter the number of pages of each item accompanying this form:

Continuation sheets of this form	0
Description	57
Claim(s)	11 DL
Abstract	0
Drawing(s)	0

10. If you are also filing any of the following, state how many against each item.

Priority documents	0
Translations of priority documents	0
Statement of inventorship and right to grant of a patent (Patents Form 7/77)	0
Request for a preliminary examination and search (Patents Form 9/77)	0
Request for a substantive examination (Patents Form 10/77)	0
Any other documents (please specify)	0

11. I/We request the grant of a patent on the basis of this application.

Signature(s)

Alan J. Walls

Date 22/08/03

12. Name, daytime telephone number and e-mail address, if any, of person to contact in the United Kingdom

ALAN J. WALLS
01737 813849

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Notes

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- Part 7 should only be completed when a divisional application is being made under section 15(4), or when an application is being made under section 8(3), 12(6) or 37(4) following an entitlement dispute. By completing part 7 you are requesting that this application takes the same filing date as an earlier UK application. If you want the new application to have the same priority date(s) as the earlier UK application, you should also complete part 6 with the priority details.

Metalloproteinase Inhibitors

The present invention relates to therapeutically active hydroxamic acid derivatives, to processes for their preparation, to pharmaceutical compositions containing them, and to the use of such compounds in medicine. In particular, the compounds are inhibitors of matrix metalloproteinases.

Background to the Invention

The matrix metalloproteinases (MMP's) are a family of zinc containing endopeptidases which are capable of cleaving large biomolecules such as the collagens, proteoglycans and gelatins. Imbalance between active MMPs and endogenous inhibitors, leads to excessive tissue disruption. The three main groups of MMPs are the collagenases, the gelatinases, and the stromelysins. Collagenases include fibroblast collagenase (MMP-1), neutrophil collagenase (MMP-8), and collagenase 3 (MMP-13). Gelatinases include 72 kDa gelatinase (gelatinase A; MMP-2) and 92 kDa gelatinase (gelatinase B; MMP-9). Stromelysins include stromelysin 1 (MMP-3), stromelysin 2 (MMP-10) and matrilysin (MMP-7). However there are MMPs which do not fit neatly into the above groups, for example metalloelastase (MMP-12), membrane-type MMP (MT-MMP or MMP-14) and stromelysin 3 (MMP-11).

Over-expression and activation of MMPs have been linked with a wide range of diseases such as cancer; rheumatoid arthritis; osteoarthritis; chronic inflammatory disorders, such as asthma, bronchitis and emphysema;

cardiovascular disorders, such as atherosclerosis; corneal ulceration; dental diseases such as gingivitis and periodontal disease; neurological disorders, such as multiple sclerosis and restenosis. For example, MMP-12 is required for the development of cigarette smoke-induced emphysema in mice, Science, 277, 2002 (1997). Inhibition of MMPs is therefore a strategy for treatment of such disease states. However, there is evidence that non-selective inhibition of matrix metalloproteinase activity may affect normal physiological process leading to dose limiting side effects. Selective inhibition of MMP-12 and/or MMP-9 is thought to be a particularly relevant strategy for intervention in inflammatory conditions.

Some MMPs can hydrolyse the membrane-bound precursor of the pro-inflammatory cytokine tumour necrosis factor α (TNF- α). This cleavage yields mature soluble TNF- α and some inhibitors of MMPs can block production of TNF- α both in vitro and in vivo. This pharmacological action is a probable contributor to the anti-inflammatory action of this class of compounds.

For a recent review of MMP inhibition as reflected in the patent literature, see Doherty et. al. Therapeutic Developments in Matrix Metalloproteinase Inhibition; Expert Opinions on Therapeutic Patents, 2002, 12, 665-707.

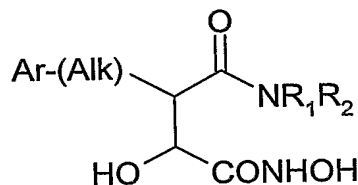
Very many of the MMP inhibitors of the prior art have a hydroxamic acid metal binding group (-CONHOH).

Brief Description of the Invention

The present invention provides a class of compounds which in general are selective inhibitors of MMP-12 relative to the collagenases and stromelysins. In addition, compounds of the invention can exhibit selective activity towards MMP-9. Compounds of the invention are therefore indicated for treatment of diseases primarily mediated by MMP-12 and/or MMP-9, especially inflammatory conditions such as multiple sclerosis and fibrosis.

Detailed Description of the Invention

According to the invention, there is provided a compound of formula (I), or an enantiomer or diastereoisomer thereof, or a salt, hydrate or solvate thereof:



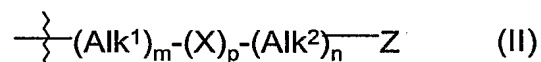
(I)

wherein

Ar represents an optionally substituted aryl, heteroaryl, C₃-C₈ cycloalkyl or heterocycloalkyl group;

Alk represents a divalent C₁-C₅ alkylene or C₂-C₅ alkenylene radical; and

R₁ and R₂ taken together with the nitrogen atom to which they are attached form a first heterocycloalkyl ring which is optionally fused to a second C₃-C₈ cycloalkyl or heterocycloalkyl ring, the said first and second rings being optionally substituted by a at least one group of formula (II):



wherein

m, p and n are independently 0 or 1;

Z represents, hydrogen, or an optionally substituted carbocyclic or heterocyclic ring of from 5 to 7 ring atoms which is optionally fused to another optionally substituted carbocyclic or heterocyclic ring of from 5 to 7 ring atoms;

Alk¹ and Alk² independently represent optionally substituted divalent C₁-C₃ alkylene radicals;

X represents -O-, -S-, -S(O)-, -S(O₂)-, -C(=O)-, -NH-, -NR₃-, -S(O₂)NH-, -S(O₂)NR₃-, -NHS(O₂)-, or -NR₃S(O₂)-, where R₃ is C₁-C₃ alkyl.

As used herein the term "(C_a-C_b)alkyl" where a and b are integers refers to a straight or branched chain alkyl moiety having from a to b carbon atoms, including for example, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, n-pentyl and n-hexyl, depending on the values of a and b.

As used herein the term "divalent (C_a-C_b)alkylene radical" where a and b are integers refers to a saturated hydrocarbon chain having from a to b carbon atoms and two unsatisfied valences.

As used herein the term "(C_a-C_b)alkenyl" where a and b are integers refers to a straight or branched chain alkenyl moiety having from a to b carbon atoms

having at least one double bond of either E or Z stereochemistry where applicable. The term includes, for example, vinyl, allyl, 1- and 2-butenyl and 2-methyl-2-propenyl, depending on the values of a and b.

As used herein the term "divalent (C_a-C_b)alkenylene radical" refers to a hydrocarbon chain having from a to b carbon atoms, at least one double bond, and two unsatisfied valences.

As used herein the unqualified term "carbocyclyl" or "carbocyclic" refers to a ring or ring system of from 3 to 14 ring atoms which are all carbon, and includes "aryl", "cycloalkyl", and "cycloalkenyl" as defined below.

As used herein the unqualified term "cycloalkyl" refers to a saturated alicyclic moiety having from 3-8 carbon atoms consisting of a single ring (e.g. cyclohexyl) or multiple condensed rings (e.g. norbornyl). Preferred cycloalkyl include cyclopentyl, cyclohexyl, norbornyl and the like.

As used herein the unqualified term "cycloalkenyl" refers to an unsaturated alicyclic moiety having from 3-8 carbon atoms and includes, for example, cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl, cycloheptenyl and cyclooctenyl. In the case of cycloalkenyl rings of from 5-8 carbon atoms, the ring may contain more than one double bond.

As used herein the unqualified term "aryl" refers to an aromatic carbocyclic group of from 6 to 14 carbon atoms having a single ring or multiple condensed rings and to two covalently linked monocyclic carbocyclic aromatic groups. Examples of aryl include phenyl, biphenyl and the like.

As used herein the unqualified term "heterocyclyl" or "heterocyclic" refers to a ring or ring system whose ring members include one or more hetero atoms selected from O, S, and N, and the term includes "heteroaryl" and "heterocycloalkyl" as defined below.

As used herein, the unqualified term "heterocycloalkyl" refers to a cycloalkyl group as defined above, in which up to 3 ring carbon atoms are replaced by heteroatoms selected from O, S and N. Preferred heterocycloalkyl include pyrrolidine, piperidine, piperazine, 1-methyl piperazine, morpholine, and the like.

As used herein the unqualified term "heteroaryl" refers to a monocyclic, or fused bicyclic or tricyclic aromatic ring or ring system containing one or more heteroatoms selected from O, S and N, and to groups consisting of two covalently linked monocyclic aromatic rings containing one or more such heteroatoms; and to groups consisting of a monocyclic carbocyclic aromatic group covalently linked to monocyclic aromatic ring containing one or more heteroatoms. Particular examples of heteroaromatic groups include optionally substituted pyridyl, pyrimidinyl, pyrrolyl, furyl, thienyl, thiophenyl, imidazolyl, oxazolyl, thiazolyl, isothiazolyl, pyrazolyl, 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, 1,2,4-triazinyl, 1,2,3-triazinyl, benzofuryl, [2,3-dihydro]benzofuryl, isobenzofuryl, benzothienyl, benzotriazolyl, isobenzothienyl, indolyl, isoindolyl, 3H-indolyl, benzimidazolyl, imidazo[1,2,-c]pyridyl, benzothiazolyl, benzoxazolyl, quinazolyl, quinazolinyl, phthalazinyl, quinoxalyl, cinnolyl, naphthyridinyl, pyrido[3,4-c]pyridyl, pyrido[3,2-c]pyridyl, pyrido[3,4,3-c]pyridyl, quinolyl, isoquinolyl, tetrazolyl, 5,6,7,8-tetrahydroquinolyl, 5,6,7,8-tetrahydroisoquinolyl, purinyl, pteridinyl, carbazolyl, xanthenyl or benzoquinolyl.

Unless otherwise specified in the context in which it occurs, the term "substituted" as applied to any moiety herein means substituted, for example, with at least one substituent selected from (C₁-C₆)alkyl, (C₁-C₆)alkoxy, hydroxy, hydroxy(C₁-C₆)alkyl, mercapto, mercapto(C₁-C₆)alkyl, (C₁-C₆)alkylthio, halo (including fluoro and chloro), trifluoromethyl, trifluoromethoxy, nitro, nitrile (-CN), oxo, phenyl, -COOH, -COOR^A, -COR^A, -SO₂R^A, -CONH₂, -SO₂NH₂, -CONHR^A, -SO₂NHR^A, -CONR^AR^B, -SO₂NR^AR^B, -NH₂, -NHR^A, -NR^AR^B, -OCONH₂, -OCONHR^A,

-OCONR^AR^B, -NHCOR^A, -NHCOOR^A, -NR^BCOOR^A, -NH₂SO₂OR^A, -NR^BSO₂OR^A, -NHCONH₂, -NR^ACONH₂, -NHCONHR^B, -NR^ACONHR^B, -NHCONR^AR^B, or -NR^ACONR^AR^B wherein R^A and R^B are independently a (C₁-C₆)alkyl or phenyl group.

The group Ar

Ar may be, for example, a 5- or 6- membered monocyclic aryl or heteroaryl ring, which is optionally substituted, for example in the 4- position in the case of a 6-membered ring, or in the 2- and/or 3- position in the case of a 5-membered ring, by at least one substituent selected from (C₁-C₃)alkyl, (C₁-C₃)alkoxy, hydroxy, hydroxy(C₁-C₃)alkyl, mercapto, mercapto(C₁-C₃)alkyl, (C₁-C₃)alkylthio, halo, trifluoromethyl, trifluoromethoxy, nitro, nitrile (-CN), -COOH, -COOR^A, -COR^A, -SO₂R^A, -CONH₂, -SO₂NH₂, -CONHR^A, -SO₂NHR^A, -CONR^AR^B, -SO₂NR^AR^B, -NH₂, -NHR^A, -NR^AR^B, -OCONH₂, -OCONHR^A, -OCONR^AR^B, -NHCOR^A, -NHCOOR^A, -NR^BCOOR^A, -NH₂SO₂OR^A, -NR^BSO₂OR^A, -NHCONH₂, -NR^ACONH₂, -NHCONHR^B, -NR^ACONHR^B, -NHCONR^AR^B, or -NR^ACONR^AR^B wherein R^A and R^B are independently C₁-C₃ alkyl, phenyl or a 5- or 6-membered monocyclic aryl or heteroaryl ring.

Ar may be, for example phenyl, 2-, 3-, or 4-pyridyl, 2-, or 3-thienyl, or 2-, or 3-furanyl, optionally substituted as specified above in relation to formula (I). Preferably the substituent can be, for example, methoxy, ethoxy, trifluoromethoxy, methyl, ethyl, trifluoromethyl, hydroxyl, mercapto, fluoro, chloro, or bromo. Presently it is preferred that Ar be 4-(C₁-C₃alkoxy)phenyl, and the most preferred being the ethoxyphenyl.

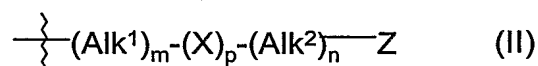
The Alk radical

Alk may be, for example, -CH₂-, -CH₂CH₂-, -CH₂CH(CH₃)-, -CH₂CH₂CH₂-, -CH₂CH₂CH₂CH₂-, -CH=CH-, -CH₂CH=CH-, -CH₂CH=CHCH₂-, or -CH=CHCH=CH-. Presently it is preferred that Alk be -CH₂CH₂CH₂-.

The group -NR₁R₂

This group is a saturated heterocyclic first ring of 5 to 7 atoms which is optionally fused to a saturated or unsaturated carbocyclic or heterocyclic second ring of 5 to 7 atoms. (The said first and/or second ring may optionally be substituted by a group of formula (II), discussed below). One of the heteroatoms of the group $-NR_1R_2$ is of course the nitrogen shown, and it may be the sole heteroatom in the ring system of there may be other nitrogen, oxygen or sulphur ring atoms.

The group $-NR_1R_2$ may be unsubstituted or substituted by at least one group (II):



It will be noted that since an optional substituent in either Alk^1 or Alk^2 may be oxo, a carbonyl group may be located adjacent the X element, forming for example combinations such as amide, reverse amide and carboxy linkages.

In one embodiment of the invention, R_1 and R_2 taken together with the nitrogen atom to which they are attached form a morpholyl group, optionally substituted by at least one group of formula (II).

In another embodiment of the invention, R_1 and R_2 taken together with the nitrogen atom to which they are attached form a thiomorpholyl group, optionally substituted by at least one group of formula (II).

In another embodiment of the invention, R_1 and R_2 taken together with the nitrogen atom to which they are attached form a piperidinyl group, optionally substituted by at least one group of formula (II).

In another embodiment of the invention, R_1 and R_2 taken together with the nitrogen atom to which they are attached form a pyrrolidinyl group, optionally substituted by at least one group of formula (II).

In another embodiment of the invention, R_1 and R_2 taken together with the nitrogen atom to which they are attached form a piperazinyl group, optionally substituted by at least one group of formula (II).

In one particular embodiment of the invention, group (II) is such that p is 0, Z is hydrogen and at least one of n and m is 1. In this subclass the group (II) is optionally substituted C_1 - C_6 alkyl, which may be linked to a ring carbon or to a ring nitrogen of the $-NR_1R_2$ group. For example when $-NR_1R_2$ is piperidinyl or piperazinyl, the 4-C (in the former) and the 4-N (in the latter) may be substituted by methyl, ethyl, or n - or iso-propyl.

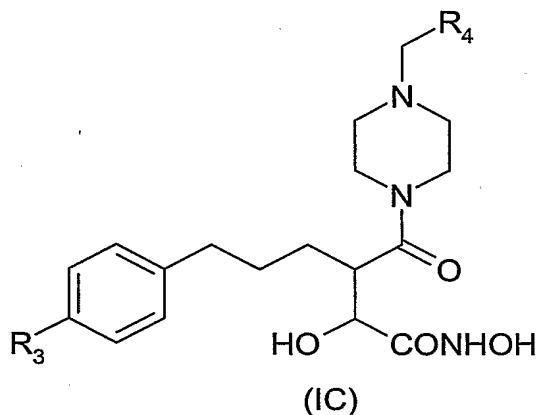
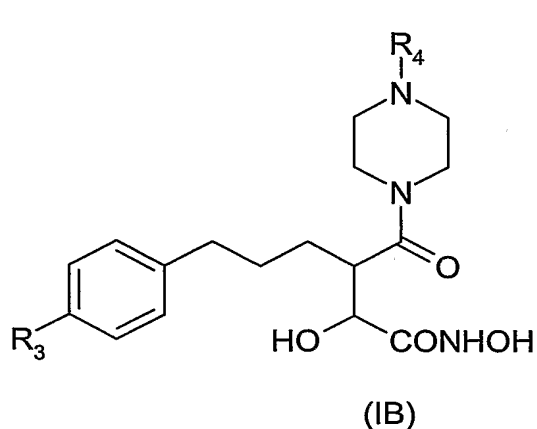
In a second particular embodiment of the invention, group (II) is such that m , n and p are all 0 and Z is a carbocyclic or heterocyclic ring directly linked to a ring carbon or ring nitrogen of the $-NR_1R_2$ group. Examples of such directly linked rings Z include cyclopentyl and cyclohexyl, and (preferably) aryl or heteroaryl rings such as phenyl, pyridyl, thienyl, furanyl, and pyrimidinyl. These directly linked rings may themselves be substituted by optional substituents, for example methoxy, ethoxy, n - or iso-propoxy, trifluoromethoxy, methylenedioxy, ethylenedioxy, methyl, ethyl, n - or isopropyl, trifluoromethyl, fluoro, chloro, bromo, methylsulfonyl, phenylsulfonyl, or mono- or di- $(C_1$ - $C_3)$ alkylamino.

In a third particular embodiment of the invention, group (II) is such that p is 0, at least one of m and n is 1, and Z is a carbocyclic or heterocyclic ring linked to a ring carbon or ring nitrogen of the $-NR_1R_2$ group via a C_1 - C_6 alkylene linker between Z and the $-NR_1R_2$ ring. In this case, the ring Z may be any of those optionally substituted Z rings discussed and preferred above in the case of the second subclass of groups (II), but here Z is linked to the $-NR_1R_2$ ring via an optionally substituted C_1 - C_6 alkylene linker radical, such as a $-CH_2-$ or $-CH_2CH_2-$ radical.

In a fourth particular embodiment of the invention, group (II) is such that p is 1, so that the group (II) contains the X heteroatom. Clearly when m is 0, X is directly linked to the $-NR_1R_2$ ring; when m and n are both 1 X interrupts a C_1 - C_6 alkylene linker between Z and the $-NR_1R_2$ ring; and when one of m and n is 1 and the other 0, the group (II) represents a variety of O, S- or N containing

substituents either directly linked to the $-NR_1R_2$ ring, or linked via a C_1 - C_3 alkylene linker.

A particular embodiment of the invention comprises compounds of formula (IB) or (IC) and salts, hydrates and solvates thereof, especially compounds having the stereoconfiguration shown in formula (IA above):



wherein R₃ is trifluoromethyl, trifluoromethoxy C_1 - C_3 alkoxy, hydroxy, or halo; R₄ is (i) $-SO_2R_5$ or $-COR_5$ wherein R₅ is C_1 - C_6 alkyl or phenyl or monocyclic heteroaryl having 5 or 6 ring atoms, optionally substituted by (C_1 - C_3)alkyl, (C_1 - C_3)alkoxy, hydroxy, hydroxy(C_1 - C_3)alkyl, mercapto, mercapto(C_1 - C_3)alkyl, (C_1 - C_3)alkylthio, halo, trifluoromethyl, trifluoromethoxy or (ii) phenyl or monocyclic heteroaryl having 5 or 6 ring atoms; optionally substituted by (C_1 - C_3)alkyl, (C_1 - C_3)alkoxy, hydroxy, hydroxy(C_1 - C_3)alkyl, mercapto, mercapto(C_1 - C_3)alkyl, (C_1 - C_3)alkylthio, halo, trifluoromethyl, trifluoromethoxy). Examples of heteroaryl rings forming part of R₄ in this embodiment include pyridyl, pyrimidinyl, triazinyl, thienyl, and furanyl.

Specific embodiments of the invention are compounds selected from the group consisting of the following:

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(pyrrolidine-1-carbonyl)-hexanoic acid hydroxyamide.

3R-(6,7-dimethoxy-3,4-dihydro-1H-isoquinoline-2-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(3-methoxy-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-methoxy-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-4-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(2RS-methyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(2,6-RS-dimethyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(thiomorpholine-4-carbonyl)-hexanoic acid hydroxyamide.

3R-(4-benzyl-piperidine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-(4-benzo[1,3]dioxol-5-ylmethyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-4-ylmethyl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-benzylpiperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyrimidin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-trifluoromethyl-pyrimidin-2-yl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-chloro-pyrimidin-2-yl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

3R-[4-(4,6-dimethoxy-[1,3,5]triazin-2-yl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(3-trifluoromethyl-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

3R-[4-(acetyl-methyl-amino)-piperidine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(methyl-propyl-amino)-piperidine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-benzyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-isobutyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-phenyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

3R-(4-benzyl-3RS-methyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-(3S,4-dibenzyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-(4-benzyl-3RS-phenyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

4-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-2S,N-dihydroxy-4-oxo-3R-(4-trifluoromethoxy-benzyl)-butyramide.

3R-benzyl-2S,N-dihydroxy-4-morpholin-4-yl-4-oxo-butyramide.

3R-(4-Benzyloxy-benzyl)-2S,N-dihydroxy-4-oxo-4-piperidin-1-yl-butyramide.

2S,N-dihydroxy-3R-(4-hydroxy-benzyl)-4-oxo-4-piperidin-1-yl-butyramide.

4-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-3R-(4-benzyloxy-benzyl)-2S,N-dihydroxy-4-oxo-butyramide.

6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

3R-(4-benzyl-piperidine-1-carbonyl)-6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(3,5-bis-trifluoromethyl-phenyl)-3R-(6,7-dimethoxy-3,4-dihydro-1H-isoquinoline-2-carbonyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(pyrrolidine-1-carbonyl)-hexanoic acid hydroxyamide

3R-(2S-benzyl-4-methyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(toluene-4-sulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

3R-[4-(5-bromo-thiophene-2-sulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-[4-(5-benzenesulfonyl-thiophene-2-sulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-[4-(4-butoxy-benzenesulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-methoxy-2,3,6-trimethyl-benzenesulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

3R-[4-(3,4-dimethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-methoxy-phenyl)-2S-hydroxy-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-methoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-fluoro-phenyl)-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-fluoro-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

As mentioned above, the present compounds are useful in human or veterinary medicine since they are active as inhibitors of MMPs, in particular as selective inhibitors of MMP-12 (and/or MMP-9) relative to MMP-1 and other collagenases and stromelysins. Accordingly in another aspect, this invention concerns:

(i) a method of management (by which is meant treatment or prophylaxis) of diseases or conditions responsive to inhibition of MMP-12 and/or MMP-9 in mammals, in particular in humans, which method comprises administering to the mammal an effective amount of a compound of formula (I) defined above, or a pharmaceutically acceptable salt thereof; and

(ii) a compound of formula (I) defined above, for use in human or veterinary medicine, particularly in the management (by which is meant treatment or prophylaxis) of diseases or conditions responsive to inhibition of MMP-12 and/or MMP-9. (Diseases or conditions responsive to inhibition of MMP-12 and/or MMP-9 include bone resorption, tumour growth or invasion by secondary metastases; rheumatoid arthritis, septic arthritis, osteoarthritis, periodontitis, gingivitis, corneal ulceration, cardiac hypertrophy, acute respiratory distress syndrome, neuroinflammatory disorders, e.g. multiple

sclerosis; restenosis; emphysema; fibrotic disease e.g. fibrosis post radiotherapy, keratin scarring, liver fibrosis and cystic fibrosis; chronic obstructive pulmonary disease; bronchitis; asthma; autoimmune disease; transplant rejection (e.g. graft versus host disease); cystic fibrosis; psoriasis; psoriatic arthritis; degenerative cartilage loss; inflammatory gastric conditions, e.g. Crohn's disease, inflammatory bowel disease, and ulcerative colitis; atopic dermatitis, epidermolysis bullosa; epidermal ulceration; a neuropathy or nephropathy e.g. interstitial nephritis, glomerulonephritis or renal failure; ocular inflammation; liver cirrhosis, Sjögren's syndrome; or an inflammatory condition of the nervous system.); and

(iii) a compound of formula (I) for use as a medicament; and

(iv) the use of a compound of formula (I) defined above in the preparation of an agent for the management (by which is meant treatment or prophylaxis) of diseases or conditions responsive to inhibition of MMP-12 and/or MMP-9; and

(v) the use of a compound of formula (I) for the preparation of a medicament for the prevention or treatment of a disease selected from inflammatory diseases including multiple sclerosis, emphysema, bronchitis, asthma, and a disease related to MMP-12 and/or MMP-9.

In a further aspect of the invention there is provided a pharmaceutical or veterinary composition comprising a compound of formula (I) defined above together with a pharmaceutically or veterinarily acceptable excipient or carrier.

It will be understood that the specific dose level for any particular patient will depend upon a variety of factors including the activity of the specific compound employed, the age, body weight, general health, sex, diet, time of administration, route of administration, rate of excretion, drug combination and the severity of the particular disease undergoing therapy. Optimum dose levels and frequency of dosing will be determined by clinical trial.

The compounds with which the invention is concerned may be prepared for administration by any route consistent with their pharmacokinetic properties. The orally administrable compositions may be in the form of tablets, capsules, powders, granules, lozenges, liquid or gel preparations, such as oral, topical, or sterile parenteral solutions or suspensions. Tablets and capsules for oral administration may be in unit dose presentation form, and may contain conventional excipients such as binding agents, for example syrup, acacia, gelatin, sorbitol, tragacanth, or polyvinyl-pyrrolidone; fillers for example lactose, sugar, maize-starch, calcium phosphate, sorbitol or glycine; tableting lubricant, for example magnesium stearate, talc, polyethylene glycol or silica; disintegrants for example potato starch, or acceptable wetting agents such as sodium lauryl sulphate. The tablets may be coated according to methods well known in normal pharmaceutical practice. Oral liquid preparations may be in the form of, for example, aqueous or oily suspensions, solutions, emulsions, syrups or elixirs, or may be presented as a dry product for reconstitution with water or other suitable vehicle before use. Such liquid preparations may contain conventional additives such as suspending agents, for example sorbitol, syrup, methyl cellulose, glucose syrup, gelatin hydrogenated edible fats; emulsifying agents, for example lecithin, sorbitan monooleate, or acacia; non-aqueous vehicles (which may include edible oils), for example almond oil, fractionated coconut oil, oily esters such as glycerine, propylene glycol, or ethyl alcohol; preservatives, for example methyl or propyl p-hydroxybenzoate or sorbic acid, and if desired conventional flavouring or colouring agents.

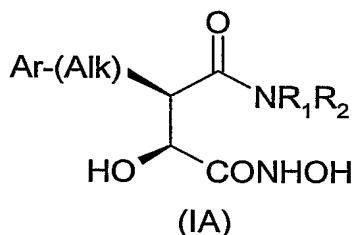
For topical application to the skin, the drug may be made up into a cream, lotion or ointment. Cream or ointment formulations which may be used for the drug are conventional formulations well known in the art, for example as described in standard textbooks of pharmaceuticals such as the British Pharmacopoeia.

For topical application to the eye, the drug may be made up into a solution or suspension in a suitable sterile aqueous or non aqueous vehicle. Additives, for instance buffers such as sodium metabisulphite or disodium edeate; preservatives including bactericidal and fungicidal agents such as phenyl

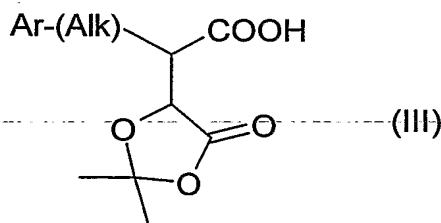
mercuric acetate or nitrate, benzalkonium chloride or chlorhexidine, and thickening agents such as hypromellose may also be included.

The active ingredient may also be administered parenterally in a sterile medium. Depending on the vehicle and concentration used, the drug can either be suspended or dissolved in the vehicle. Advantageously, adjuvants such as a local anaesthetic, preservative and buffering agents can be dissolved in the vehicle.

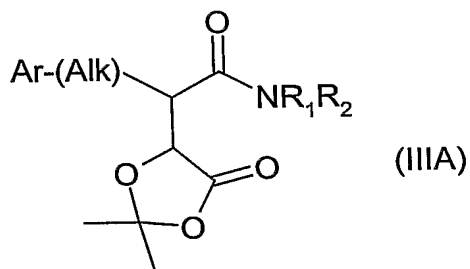
In one embodiment of the invention, compounds of the invention have the stereochemical configuration shown in formula (IA):



Compounds of formula (I) may be prepared by standard literature methods, as illustrated in the Examples herein. In general, the compounds may be prepared by coupling a dioxolane-protected di-carboxylic acid of formula (III) with the desired cyclic amine HNR_1R_2 :



wherein R_1 , R_2 : Ar and Alk are as defined in relation to formula (I), to form the intermediate (IIIA)



then reacting intermediate (IIIA) with hydroxylamine.

Methods of coupling amines with carboxylic acids to form amides are very well known, for example from the art of peptide synthesis.

Thus, the invention also includes

(vi) a compound of formula (IIIB) wherein R_1 , R_2 , Ar and Alk are as defined in relation to formula (I)

(vii) a process for the preparation of a compound of formula (I) defined above, comprising the deprotection and/or transformation of a compound of formula (IIIA) as defined above,) wherein R_1 , R_2 , Ar and Alk are as defined in relation to formula (I)

(viii) a process for the preparation of a compound of formula (IIIA) comprising the step of reacting a compound of formula (III) with a cyclic amine HNR_1R_2 , wherein R_1 , R_2 : Ar and Alk are as defined in relation to formula (I),

The following preparative Examples describe the preparation of compounds useful in accordance with the invention

The following abbreviations have been used in the examples:

AcOEt – Ethyl acetate

CH_3CN - Acetonitrile

DMF - N,N-Dimethylformamide

HOBT - 1-Hydroxybenzotriazole

HOAT - 1-Hydroxyazobenzotriazole

MgSO₄ – Magnesium sulfate

Pfp – Pentafluorophenol

WSCDI - N-(3-Dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride

HCl - Hydrochloric acid

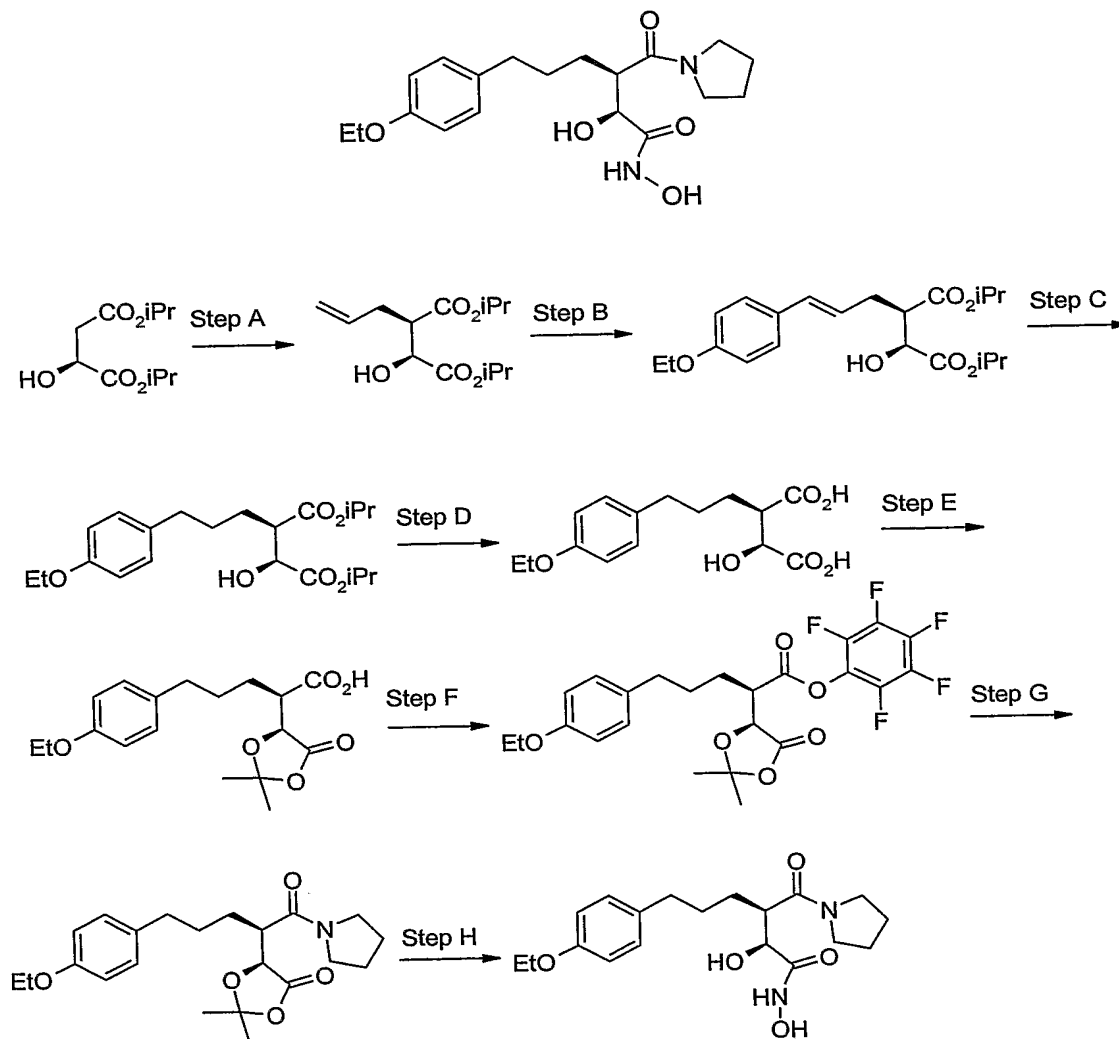
P(O-Tol)₃ – Tri-O-tolylphosphine

THF – Tetrahydrofuran

TFA – Trifluoroacetic acid

Z - Benzyloxycarbonyl

Example 1: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(pyrrolidine-1-carbonyl)-hexanoic acid hydroxyamide.



Reagents and conditions. A: LiHMDS, AlBr, THF, -78C to RT; B: ArBr, P(o-Tol)₃, Pd(OAc)₂, NEt₃, CH₃CN; C: 10%Pd/C, H₂, MeOH; D: LiOH, MeOH, H₂O; E: CuCl₂, dimethoxypropane, acetone; F: pentafluorophenol, WSCDI, HOAt, CH₂Cl₂; G: pyrrolidine, NEt₃, CH₂Cl₂; H: aq NH₂OH, iPrOH

Scheme 1

Example 1 was prepared as outlined in Scheme 1 using procedures described below.

Step A: 2R-allyl-3S-hydroxy-succinic acid diisopropylester.

To a cold (-78C) solution of 2S-hydroxy-succinic acid diisopropyl ester (19.70 ml, 95 mmol) in THF (35 ml) was added LiHMDS (200 ml, 0.2 mol, 2.1 eq.)

dropwise. The reaction mixture was stirred at -78C for two hours and then at -30C for 30 min. The reaction mixture was then cooled to -78C and allyl bromide (12.36 ml, 0.14 mol, 1.5 eq.) added dropwise. The reaction mixture was then allowed to warm to RT overnight. It was poured into a saturated solution of NH_4Cl /ice (200 ml). Extraction with AcOEt (3 X 200 ml) followed by a wash with water (50 ml) and with brine (50 ml) afforded a yellow oil after removal of the solvents under vacuo. Purification by flash chromatography gave 2R-allyl-3S-hydroxy-succinic acid diisopropylester as a colourless oil (7.76 g, de = 80%, 40% yield).

$^1\text{H-NMR}$; delta (CDCl_3): 5.77-5.88 (1H, m), 4.98-5.21 (4H, m), 4.22 (1H, brs), 3.18 (1H, brs), 2.87-2.94 (1H, m), 2.56-2.65 (1H, m), 2.40-2.48 (1H, m), 1.29 (6H, d, $J=6.3$ Hz), 1.22 (6H, d, $J=6.3$ Hz).

LRMS: +ve ion 281 ($\text{M}+\text{Na}$).

Step B: 2R-[3-(4-ethoxy-phenyl)-allyl]-3S-hydroxy-succinic acid diisopropyl ester.

To a solution of 2R-allyl-3S-hydroxy-succinic acid diisopropylester (4.79 g, 18.5 mmol), 4-bromo phenetole (3.19 ml, 22.2 mmol, 1.2 eq.) and NEt_3 (6.22 ml, 44.6 mmol, 2.4 eq.) in CH_3CN (40 ml), was added a sonicated (for 2 min) suspension of $\text{P}(\text{O-Tol})_3$ (0.57 g, 2.22 mmol, 0.1 eq.) and $\text{Pd}(\text{OAc})_2$ (209 mg, 5%) in CH_3CN (5 ml). The reaction mixture was heated to reflux for 2 hrs. CH_3CN was removed under vacuo. The crude was extracted with AcOEt (3 X 200 ml), washed with water (50 ml) and with brine (50 ml). A purification by flash chromatography afforded the desired 2R-[3-(4-ethoxy-phenyl)-allyl]-3S-hydroxy-succinic acid diisopropyl ester (5.92 g, 84% yield).

$^1\text{H-NMR}$; delta (CDCl_3): 7.28 (2H, d, $J=8.8$ Hz), 6.83 (2H, d, $J=8.8$), 6.46 (1H, d, $J=15.7$ Hz), 6.02-6.12 (1H, m), 4.98-5.13 (2H, m), 4.26 (1H, dd, $J=7.1$, 3.0 Hz), 4.02 (2H, q, $J=7.0$ Hz): 3.23 (1H, d, $J=7.1$ Hz), 2.92-2.97 (1H, m), 2.68-2.79 (1H, m), 2.49-2.62 (1H, m), 1.41 (3H, t, $J=7.0$ Hz), 1.19-1.30 (12H, m).

LRMS: +ve ion 401 ($\text{M}+\text{Na}$).

Step C: 2R-[3-(4-ethoxy-phenyl)-propyl]-3S-hydroxy-succinic acid diisopropyl ester.

To a solution of 2R-[3-(4-ethoxy-phenyl)-allyl]-3S-hydroxy-succinic acid diisopropyl ester (129 mg, 0.34 mmol) in MeOH (10 ml) under an inert atmosphere, was added 10%Pd/C (13 mg). H₂ was bubbled through the resulting suspension for 30 min. The reaction mixture was then stirred under 1 atmosphere of H₂ for 16 hrs. Pd/C was filtered off and the solvent removed under reduced pressure to give 2R-[3-(4-ethoxy-phenyl)-propyl]-3S-hydroxy-succinic acid diisopropyl ester (115 mg, 88% yield).

¹H-NMR; delta (CDCl₃): 7.08 (2H, d, J=8.6 Hz), 6.81 (2H, d, J=8.6), 4.97-5.14 (2H, m), 4.20 (1H, dd, J=7.3, 3.5 Hz), 4.01 (2H, q, J=7.0 Hz), 3.18 (1H, d, J=7.3 Hz), 2.77-2.83 (1H, m), 2.55-2.62 (2H, m), 1.45-1.94 (4H, m), 1.40 (3H, t, J=7.0 Hz), 1.12-1.30 (12H, m).

LRMS: +ve ion xx (M+Na).

Step D: 2R-[3-(4-ethoxy-phenyl)-propyl]-3S-hydroxy-succinic acid.

To a solution of 2R-[3-(4-ethoxy-phenyl)-propyl]-3S-hydroxy-succinic acid diisopropyl ester (4.78 g, 12.6 mmol) in THF/water (3:1, 120 ml) was added NaOH (1.66 g, 41.5 mmol, 5.5 eq.). The reaction mixture was then stirred for 16 hrs at RT. The mixture was concentrated under reduced pressure and acidify to pH = 3 by addition of HCl 1N. The hydroxy diacid was extracted with AcOEt. The organic layer was dried over MgSO₄ and the solvent removed under reduced pressure to give the desired 2R-[3-(4-ethoxy-phenyl)-propyl]-3S-hydroxy-succinic acid (3.66 g, 85% yield).

¹H-NMR; delta (MeOD): 7.07 (2H, d, J=8.6 Hz), 6.79 (2H, d, J=8.6), 4.23 (1H, d, J=5.8 Hz), 3.98 (2H, q, J=7.0 Hz), 2.76-2.81 (1H, m), 2.53-2.59 (2H, m), 1.55-1.72 (4H, m), 1.35 (3H, t, J=7.0 Hz).

LRMS: +ve ion 319 (M+Na); -ve ion 295 (M-H).

Step E: 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid.

To a solution of 2R-[3-(4-ethoxy-phenyl)-propyl]-3S-hydroxy-succinic acid (3.66 g, 12.3 mmol) in acetone (50 ml) under an inert atmosphere were added dimethoxy propane (2.58 ml, 21 mmol, 1.7 eq.) and copper chloride (165 mg, 1.2 mmol, 0.1 eq.). The reaction mixture was stirred at RT for 16 hrs. The solvent was then removed under vacuo to give 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid (4.03 g, 97% yield).

¹H-NMR; delta (CDCl₃): 7.08 (2H, d, J=8.5 Hz), 6.82 (2H, d, J=8.5), 4.48 (1H, d, J=4.8 Hz), 4.01 (2H, q, J=7.0 Hz), 2.91-2.98 (1H, m), 2.54-2.64 (3H, m), 1.23-2.20 (4H, m), 1.58 (3H, s), 1.53 (3H, s), 1.40 (3H, t, J=7.0 Hz).

LRMS: +ve ion 359 (M+Na); -ve ion 335 (M-H).

Step F. 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid pentafluorophenyl ester.

To a cold (0C) solution of 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid (4.03g, 12 mmol) and pentafluoro phenol (2.43 g, 13.2 mmol, 1.1 eq.) in CH₂Cl₂ (50 ml) was added WSC (2.54 g, 13.2 mmol, 1.1 eq.). The reaction mixture was allowed to warm to RT overnight. CH₂Cl₂ was removed under vacuo and the resulting crude reaction mixture was dissolved in AcOEt (200 ml). The organic layer was washed with water (50 ml), NaHCO₃ sat (20 ml) and finally with brine (20 ml). Solvent was removed under reduced pressure to give an oil which was purified by flash chromatography to furnish the expected 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid pentafluorophenyl ester (3.94 g, 65% yield).

¹H-NMR; delta (CDCl₃): 7.09 (2H, d, J=8.4 Hz), 6.83 (2H, d, J=8.4 Hz), 4.56 (1H, d, J=6.0 Hz), 4.01 (2H, q, J=7.0 Hz), 3.20-3.28 (1H, m), 2.64 (2H, t, J=7.6 Hz), 1.98-2.08 (2H, m), 1.70-1.86 (2H, m), 1.62 (3H, s), 1.57 (3H, s), 1.40 (3H, t, J= 7.0 Hz).

Step G. 5-(4-Ethoxy-phenyl)-2R-[(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)]-1-pyrrolidin-1-yl-pentan-1-one.

To a solution of 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid pentafluorophenyl ester (150 mg, 0.30 mmol) in CH_2Cl_2 (10 ml) was added pyrrolidine (30 μl , 0.36 mmol, 1.2 eq.). The reaction mixture was stirred for 16 hrs and the solvent was removed under vacuo. The crude was taken-up in AcOEt (70 ml) and washed with water (10 ml), then with NaHCO_3 sat (10 ml) and finally with brine (10 ml). The solvent was dried over MgSO_4 and removed under reduced pressure to give the desired 5-(4-Ethoxy-phenyl)-2R-[(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)]-1-pyrrolidin-1-yl-pentan-1-one (116 mg, quant.).

$^1\text{H-NMR}$; delta (CDCl_3): 7.05 (2H, d, $J=8.6\text{Hz}$), 6.8 (2H, d, $J=8.6\text{Hz}$), 4.55 (1H, d, $J=8.4\text{Hz}$), 3.99 (2H, m), 3.8-3.3 (10H, m), 3.05 (1H, m), 2.55 (2H, t, $J=7.6\text{Hz}$), 2.1-1.7 (2H, m), 1.6 (3H, s), 1.5 (3H, s), 1.4 (3H, t, $J=7.0\text{Hz}$).

LRMS: +ve ion 405 (M+H), 428 (M+Na).

Step H. 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(pyrrolidine-1-carbonyl)-hexanoic acid hydroxyamide

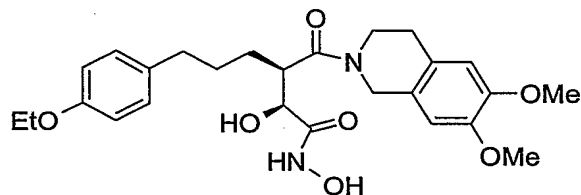
To a solution of 5-(4-Ethoxy-phenyl)-2R-[(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)]-1-pyrrolidin-1-yl-pentan-1-one (116 mg, 0.30 mmol) in *i*-PrOH (5 ml), was added an aqueous solution of hydroxylamine (50%, 99 μl , 1.5 mmol, 5 eq.). The reaction mixture was allowed to stir at RT for 16 hrs. The solvent was removed under reduced pressure to yield an oil which was purified by preparative reverse phase chromatography to give the required product.

$^1\text{H-NMR}$; delta (CD_3OD): 7.05 (2H, d, $J=8.6\text{Hz}$), 6.8 (2H, d, $J=8.6\text{Hz}$), 4.0 (4H, m), 3.85 (1H, m), 3.7 (1H, m), 3.4 (2H, m), 3.1 (1H, m), 2.55 (2H, m), 1.9-1.5 (7H, m), 1.35 (3H, t, $J=7.0\text{Hz}$).

LRMS: +ve ion 387 (M+Na); -ve ion 363 (M-H)

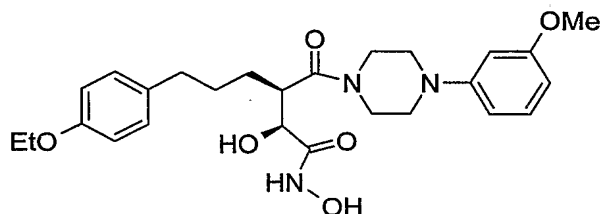
The compounds of Examples 2-20 were prepared by the method of Example 1 by parallel synthesis, using the appropriate amine in Step G. The products were purified by preparative HPLC.

Example 2: 3R-(6,7-dimethoxy-3,4-dihydro-1H-isoquinoline-2-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



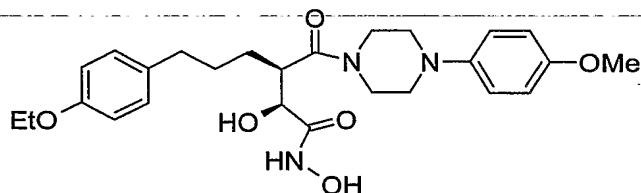
LRMS: +ve ion 487 (M+H), 509 (M+Na); -ve ion 485 (M-H).

Example 3: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(3-methoxy-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



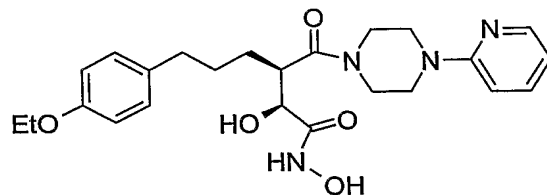
LRMS: +ve ion 486 (M+H), 508 (M+Na); -ve ion 484 (M-H).

Example 4: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-methoxy-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



LRMS: +ve ion 486 (M+H), 508 (M+Na); -ve ion 484 (M-H).

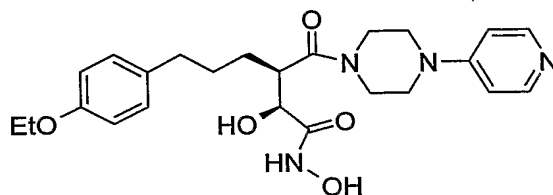
Example 5: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.



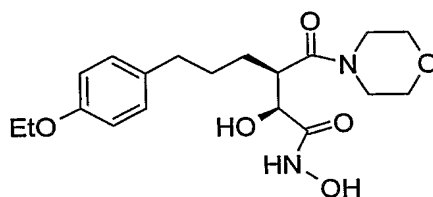
$^1\text{H-NMR}$; delta (CD_3OD): 8.1 (1H, d, $J = 1.4\text{Hz}$), 7.55 (1H, m), 7.05 (2H, d, $J = 8.7\text{Hz}$), 6.9-6.6 (4H, m), 3.95 (1H, d, $J = 7.0\text{Hz}$), 3.55 (4H, m), 2.55 (1H, m), 1.8-1.5 (6H, m), 1.35 (3H, m).

LRMS: +ve ion 457 (M+H); -ve ion 455 (M-H).

Example 6: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-4-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.



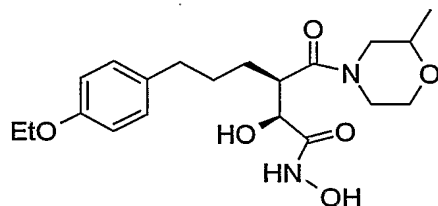
Example 7: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(morpholine-4-carbonyl)-hexanoic acid hydroxyamide.



$^1\text{H-NMR}$; delta (CD_3OD): 7.05 (2H, d, $J = 8.6\text{Hz}$), 6.8 (2H, d, $J = 8.6\text{Hz}$), 4.05-3.90 (3H, m), 3.8-3.4 (8H, m), 2.55 (2H, t, $J = 6.7\text{Hz}$), 1.75-1.4 (4H, m), 1.35 (3H, t, $J = 7.0\text{Hz}$).

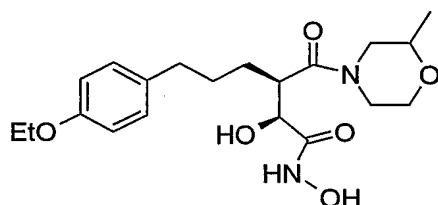
LRMS: +ve ion 403 (M+Na); -ve ion 379 (M-H).

Example 8: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(2RS-methyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.



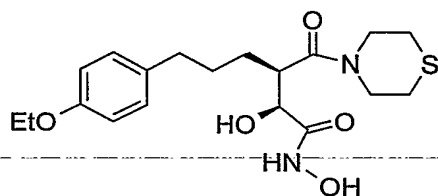
LRMS: +ve ion 417 (M+Na), 395 (M+H); -ve ion 393 (M-H).

Example 9: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(2,6-RS-dimethyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.



LRMS: +ve ion 431 (M+Na), 409 (M+H); -ve ion 407 (M-H).

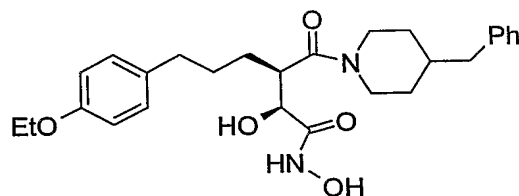
Example 10: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(thiomorpholine-4-carbonyl)-hexanoic acid hydroxyamide.



¹H-NMR; delta (CD₃OD): 7.05 (2H, d), 6.8 (2H, d), 4.0 (5H, m), 3.8-3.5 (2H, m), 2.9-2.4 (7H, m), 1.55 (4H, m) and 1.3 (3H, t).

LRMS: +ve ion 419 (M+Na). 397 (M+H); -ve ion 395 (M-H).

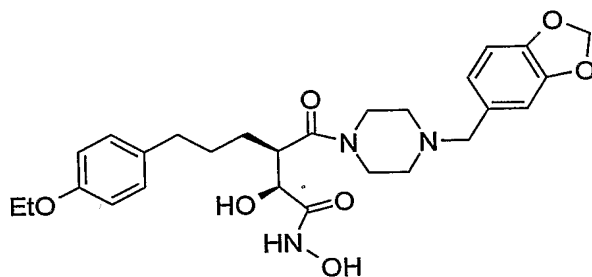
Example 11: 3R-(4-benzyl-piperidine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



$^1\text{H-NMR}$; delta (CD_3OD): 7.3-7.0 (8H, m), 6.8 (2H, m), 4.55 (1H, d, $J=12.4\text{Hz}$), 4.05 (2H, dd, $J=2.0\text{Hz}$), 3.9 (2H, m), 2.9 (1H, m), 2.6-2.4 (5H, m), 1.84 (1H, d, $J=2.9\text{Hz}$), 1.7-1.5 (6H, m), 1.35 (3H, t, $J=7.0\text{Hz}$).

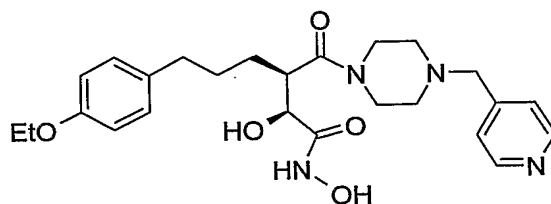
LRMS: +ve ion 491 ($\text{M}+\text{Na}$); -ve ion 467 ($\text{M}-\text{H}$).

Example 12: 3R-(4-benzo[1,3]dioxol-5-ylmethyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



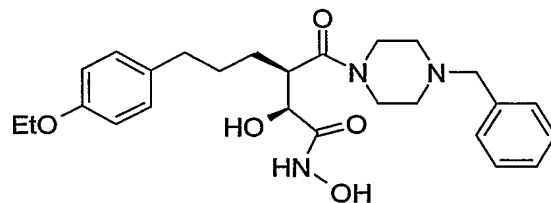
LRMS: +ve ion 514 ($\text{M}+\text{H}$); -ve ion 512 ($\text{M}-\text{H}$).

Example 13: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-4-ylmethyl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.



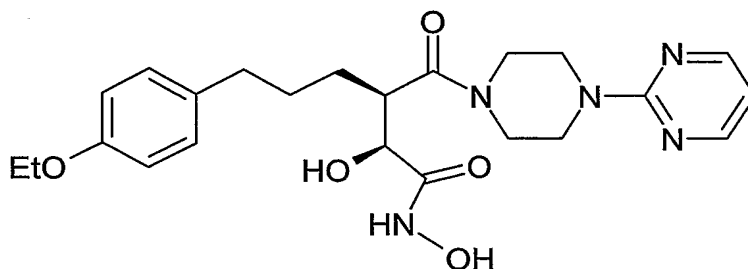
LRMS: +ve ion 471 ($\text{M}+\text{H}$); -ve ion 469 ($\text{M}-\text{H}$).

Example 14: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-benzylpiperazine-1-carbonyl)-hexanoic acid hydroxyamide.



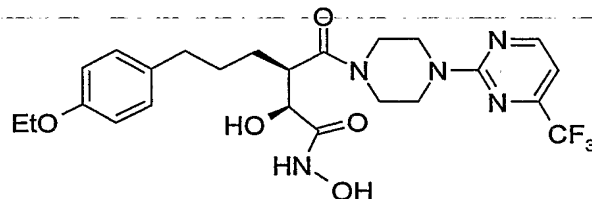
LRMS: +ve ion 471 (M+H); -ve ion 469 (M-H).

Example 15: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyrimidin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.



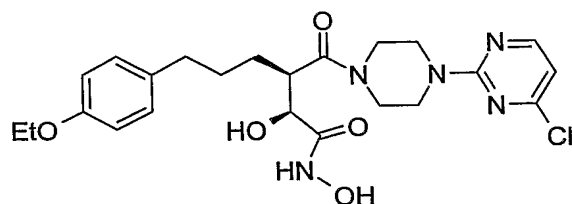
LRMS: +ve ion 458 (M+H); -ve ion 456 (M-H).

Example 16: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-trifluoromethyl-pyrimidin-2-yl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



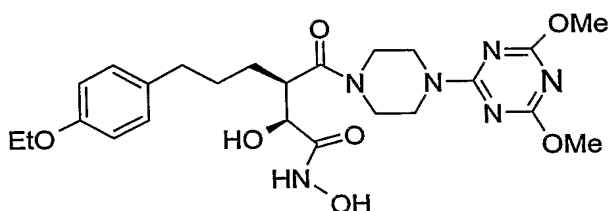
LRMS: +ve ion 526 (M+H), 548 (M+Na); -ve ion 524 (M-H).

Example 17: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-chloro-pyrimidin-2-yl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



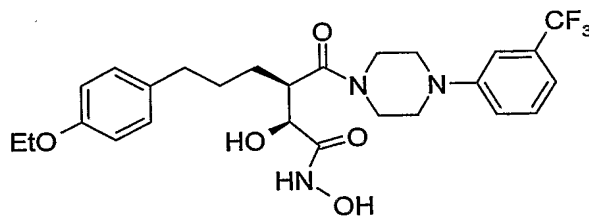
LRMS: +ve ion 492 (M+H); -ve ion 490 (M-H).

Example 18: 3R-[4-(4,6-dimethoxy-[1,3,5]triazin-2-yl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



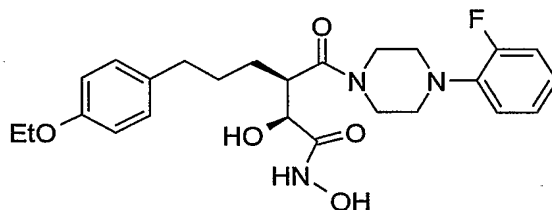
LRMS: +ve ion 519 (M+H); -ve ion 517 (M-H).

Example 19: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(3-trifluoromethyl-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



LRMS: +ve ion 524 (M+H); -ve ion 522 (M-H).

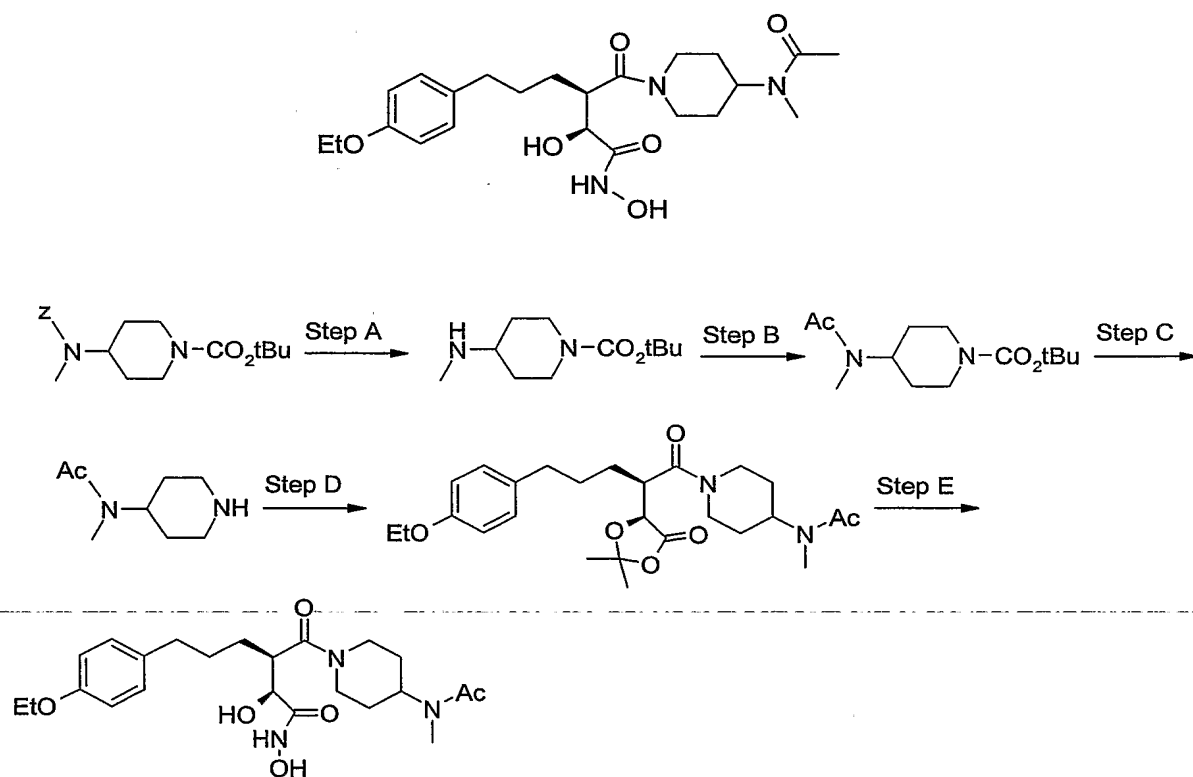
Example 20: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



LRMS: +ve ion 474 (M+H); -ve ion 472 (M-H).

Example 21: 3R-[4-(acetyl-methyl-amino)-piperidine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

The compound of Example 21 was prepared as outlined in scheme 2, using procedures described below.



Reagents and conditions: A: H_2 , Pd/C, MeOH; B: AcCl, NEt_3 , CH_2Cl_2 ; C: TFA/ CH_2Cl_2 ; D: PFP ester, NEt_3 ; E: NH_2OH , iPrOH

Scheme 2

Step A: 4-methylamino-piperidine-1-carboxylic acid tert-butyl ester.

To a solution of 4-(benzyloxycarbonyl-methyl-amino)-piperidine-1-carboxylic acid tert-butyl ester (5 g, 0.014 mmol) in MeOH (30 ml) was added under an inert atmosphere 10% Pd/C (500 mg). H₂ was bubbled through the resulting suspension for 2 hrs. The reaction mixture was then stirred under 1 atm. of H₂ for 2 hrs. Pd/C was filtered off and the solvent was removed under reduced pressure to give the expected 4-methylamino-piperidine-1-carboxylic acid tert-butyl ester (3.08 g, quant.).

Step B: 4-(acetyl-methyl-amino)-piperidine-1-carboxylic acid tert-butyl ester.

To a solution of 4-methylamino-piperidine-1-carboxylic acid tert-butyl ester (616 mg, 2.9 mmol) in CH₂Cl₂ (25 ml), was added acetyl chloride (215 μ l, 3.02 mmol, 1.05 eq.) dropwise. The reaction mixture was stirred at RT for 16 hrs. The crude reaction mixture was extracted with CH₂Cl₂ (2 X 50 ml) and washed with water (10 ml) and with brine (10 ml). The solvent was removed under reduced pressure to give the expected 4-(Acetyl-methyl-amino)-piperidine-1-carboxylic acid tert-butyl ester (708 mg, 96% yield).

¹H-NMR; delta (CDCl₃): 4.7-4.5 (1H, m), 4.1 (2H, br d, J= 12.7Hz), 2.8 (5H, m), 2.1 (5H, m), 1.6 (2H, m), 1.5 (9H, s).

LRMS: +ve ion 215 (M+H).

Step C: N-methyl-N-piperidin-4-yl-acetamide.

To a solution of 4-(acetyl-methyl-amino)-piperidine-1-carboxylic acid tert-butyl ester (525 mg, 2.05 mmol) in AcOEt (5 ml) was added HCl 3N (4 ml). The reaction mixture was stirred at RT for 16 hrs. Solvent were removed under reduced pressure to gine the expected N-methyl-N-piperidin-4-yl-acetamide as its hydrochloride salt (395 mg, quant.).

¹H-NMR; delta (CD₃OD): 4.6 (1H, m), 3.5 (2H, br d, J= 12.8Hz), 3.2-3.1 (2H, m), 3.0-2.85 (3H, s), 2.2 (3H, s), 2.1-1.8 (4H, m).

LRMS: +ve ion 157 (M+H).

Step D: N-{1-[2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoyl]-piperidin-4-yl}-N-methyl-acetamide.

To a solution of 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid pentafluorophenyl ester (230 mg, 0.46 mmol) in CH_2Cl_2 (5 ml) was added N-Methyl-N-piperidin-4-yl-acetamide (137 mg, 0.6 mmol, 1.3 eq.) followed by NEt_3 (257 μl , 1.83 mmol, 4 eq.). The reaction mixture was stirred at RT for 16 hrs. The reaction mixture was extracted with CH_2Cl_2 (2 X 20 ml) and washed with water (10 ml), NaHCO_3 sat (5 ml) and finally with brine (10 ml). The organic layer was dried over MgSO_4 and the solvent were removed under reduced pressure to give the expected N-{1-[2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoyl]-piperidin-4-yl}-N-methyl-acetamide in quantitative yield. This compound was used in the next step without any further purification.

$^1\text{H-NMR}$; delta (CDCl_3): 7.05 (2H, d, $J = 7.2\text{Hz}$), 6.7 (2H, d, $J = 7.2\text{Hz}$), 4.75 (2H, m), 4.55 (1H, d, $J = 7.9\text{Hz}$), 4.0 (3H, m), 3.05 (1H, m), 2.8 (1H, m), 2.7-2.5 (4H, m), 2.1 (3H, s), 1.85 (3H, m), 1.75-1.5 (4H, m), 1.65 (3H, s), 1.55 (3H, s), 1.4 (3H, t, $J = 7.0\text{Hz}$), 0.8 (3H, t, $J = 7.2\text{Hz}$).

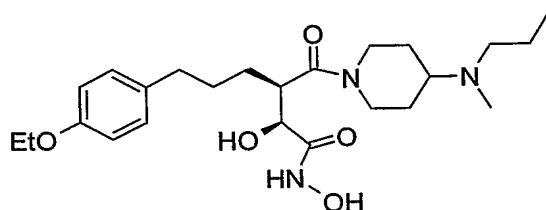
LRMS: +ve ion 475 (M+H).

Step E: 3R-[4-(acetyl-methyl-amino)-piperidine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

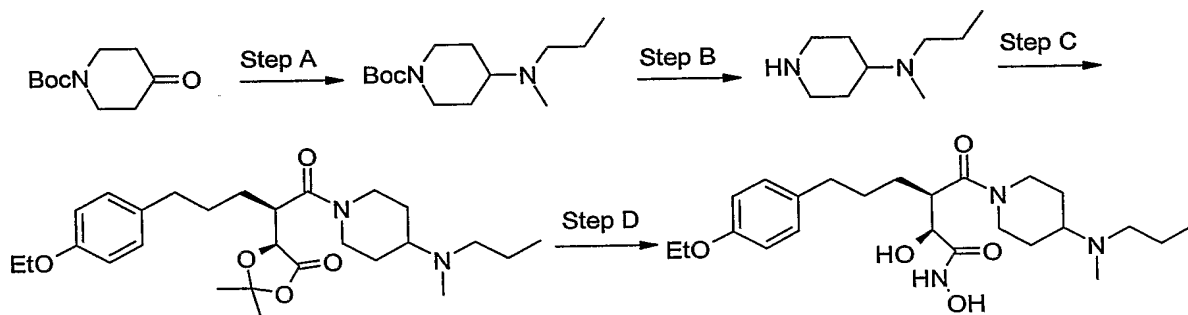
To a solution of N-{1-[2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoyl]-piperidin-4-yl}-N-methyl-acetamide (135 mg, 0.28 mmol) in *i*-PrOH (5 ml) was added an aqueous solution of hydroxylamine (50%, 94 μl , 1.43 mmol, 5 eq.). The reaction mixture was allowed to stir at RT for 16 hrs. The solvent was removed under reduced pressure to yield an oil which was purified by preparative reverse phase chromatography to give the required 3R-[4-(acetyl-methyl-amino)-piperidine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

LRMS: +ve ion.(M+Na); -ve ion (M-H)

Example 22: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(methyl-propyl-amino)-piperidine-1-carbonyl]-hexanoic acid hydroxyamide.



Example 22 was prepared as outlined in Scheme 3 using procedures described below.



Reagents and conditions: A: methyl propyl amine, ; B: TFA/CH₂Cl₂; C: PFP ester, NEt₃; D: NH₂OH, iPrOH

Scheme 3

Step A: 4-(methyl-propyl-amino)-piperidine-1-carboxylic acid tert-butyl ester.

To a solution of 4-oxo-piperidine-1-carboxylic acid tert-butyl ester (100 mg, 0.5 mmol) in MeOH (5 ml) was added NaBH₃CN (32 mg, 0.5 mmol, 1 eq.). The pH of the reaction mixture was adjusted to 5.5 by addition of 5N HCl and stirred for 48 hrs at RT under an inert atmosphere. The solvent was removed under reduced pressure and the crude was taken-up in AcOEt (150 ml). The organic layer was washed with NaHCO₃ (10 ml) and with brine (10 ml) and was dried over MgSO₄. The solvent was removed under vacuo to give the expected 4-(methyl-propyl-amino)-piperidine-1-carboxylic acid tert-butyl ester (106 mg, 82% yield).

$^1\text{H-NMR}$; delta (CDCl_3): 4.14 (1H, br d, $J = 12.2\text{Hz}$), 3.8 (1H, m), 3.0 (1H, m), 2.8-2.35 (5H, m), 2.2 (3H, s), 1.9-1.7 (3H, m), 1.5 (11H, s), 0.9 (3H, t, $J = 3.4\text{Hz}$).

LRMS: +ve ion 257 (M+H).

Step B: methyl-piperidin-4-yl-propyl-amine.

To a solution of 4-(methyl-propyl-amino)-piperidine-1-carboxylic acid tert-butyl ester (106 mg, 0.41 mmol) in AcOEt (10 ml) was added 3N HCl (4 ml). The reaction mixture was stirred for 16 hrs at RT. The solvent was removed under reduced pressure to give the desired methyl-piperidin-4-yl-propyl-amine (87 mg, 92% yield).

$^1\text{H-NMR}$; delta (CD_3OD): 3.95 (1H, m), 3.6 (2H, d, $J = 13.9\text{Hz}$), 2.3 (2H, d, $J = 11.3\text{Hz}$), 2.05 (4H, m), 1.85 (4H, m), 1.05 (3H, t, $J = 7.3\text{Hz}$).

LRMS: +ve ion 157 (M+H).

Step C: 5S-{4-(4-Ethoxy-phenyl)-1R-[4-(methyl-propyl-amino)-piperidine-1-carbonyl]-butyl}-2,2-dimethyl-[1,3]dioxolan-4-one.

To a solution of 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid pentafluorophenyl ester (175 mg, 0.35 mmol) in CH_2Cl_2 (5 ml) was added methyl-piperidin-4-yl-propyl-amine (87 mg, 0.45 mmol, 1.3 eq.) followed by NEt_3 (197 μl , 1.39 mmol, 4 eq.). The reaction mixture was stirred for 16 hrs and the solvent was removed under vacuo. The crude was taken-up in AcOEt (70 ml) and washed with water (10 ml), then with NaHCO_3 sat (10 ml) and finally with brine (10 ml). The organic layer was dried over MgSO_4 and removed under reduced pressure. A purification on silica gel gave the desired 5S-{4-(4-Ethoxy-phenyl)-1R-[4-(methyl-propyl-amino)-piperidine-1-carbonyl]-butyl}-2,2-dimethyl-[1,3]dioxolan-4-one. (82 mg, 50% yield).

$^1\text{H-NMR}$; delta (CDCl_3): 7.0 (2H, d, $J = 8.3\text{Hz}$), 6.7 (2H, d, $J = 8.3\text{Hz}$), 4.65 (1H, m), 4.5 (1H, m), 3.9 (2H, m), 3.05 (1H, m), 2.9 (1H, m), 2.5 (4H, d, $J = 7.7\text{Hz}$),

2.3 (2H, m), 2.15 (3H, dd, $J = 4.0, 13.8\text{Hz}$), 1.7 (4H, t, $J = 7.0\text{Hz}$), 1.55 (3H, s), 1.5 (3H, s), 1.45-1.3 (7H, m), 1.3 (3H, t, $J = 7.0\text{Hz}$), 0.8 (3H, t, $J = 7.2\text{Hz}$).

LRMS: +ve ion 475 (M+H).

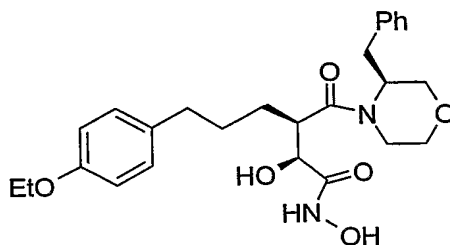
Step D: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(methyl-propyl-amino)-piperidine-1-carbonyl]-hexanoic acid hydroxyamide.

To a solution of 5S-{4-(4-ethoxy-phenyl)-1R-[4-(methyl-propyl-amino)-piperidine-1-carbonyl]-butyl}-2,2-dimethyl-[1,3]dioxolan-4-one (82 mg, 0.17 mmol) in *i*-PrOH (3 ml), was added an aqueous solution of hydroxylamine (50%, 57 μl , 0.87 mmol, 5 eq.). The reaction mixture was allowed to stir at RT for 16 hrs. The solvent was removed under reduced pressure to yield an oil which was purified by preparative reverse phase chromatography to give the required product.

$^1\text{H-NMR}$; delta (CH_3OD): 7.1 (2H, d), 6.8 (2H, d), 4.6 (1H, d), 4.1 (1H, d), 4.0 (2H, m), 3.35 (1H, m), 2.9-2.35 (8H, m), 2.3-2.2 (3H, s), 2.0-1.4 (10H, m), 1.35 (3H, t) and 0.9 (3H, t).

LRMS: +ve ion 450 (M+H); -ve ion 448 (M-H)

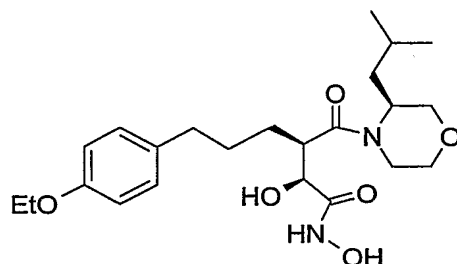
Example 23: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-benzyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.



The compound of example 23-25 were prepared by the method of example 1 according to scheme 1, and using the appropriate amine (synthesised according to the procedures described in Tetrahedron, 36, 409-415, 1980 and Journal of Organic Chemistry, 61, 4990-4998, 1996), in Step A. The products were purified by preparative HPLC.

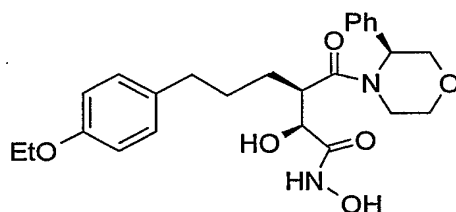
LRMS: +ve ion 471 (M+H); -ve ion 469 (M-H)

Example 24: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-isobutyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.



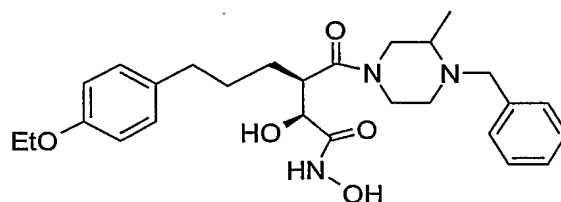
LRMS: +ve ion 437 (M+H); -ve ion 435 (M-H)

Example 25: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-phenyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.



LRMS: +ve ion 457 (M+H), 479 (M+Na); -ve ion 455 (M-H)

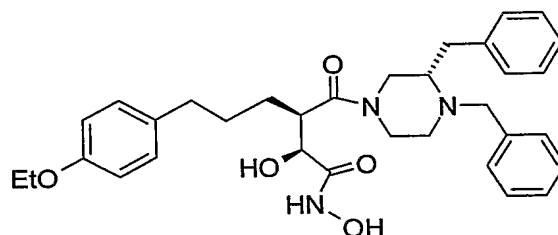
Example 26: 3R-(4-benzyl-3RS-methyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



The compound of example 26-28 were prepared by the method of example 1 according to scheme 1, and using the appropriate piperazines (synthesised according to the procedure described in *Tetrahedron: Asymmetry*, 12, 3319-3324, **2001**; and *Org. Prep. Proced. Int.*, 22, (6), 761-768, **1990**, in Step A. The products were purified by preparative HPLC.

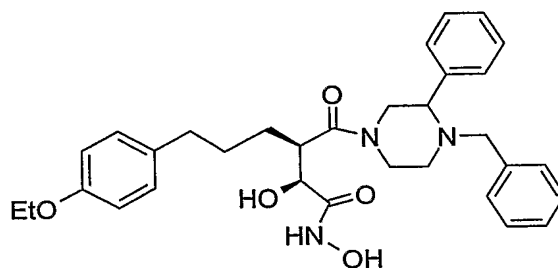
LRMS: +ve ion 484 (M+H).

Example 27: 3R-(3S,4-dibenzyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



LRMS: +ve ion 560 (M+H).

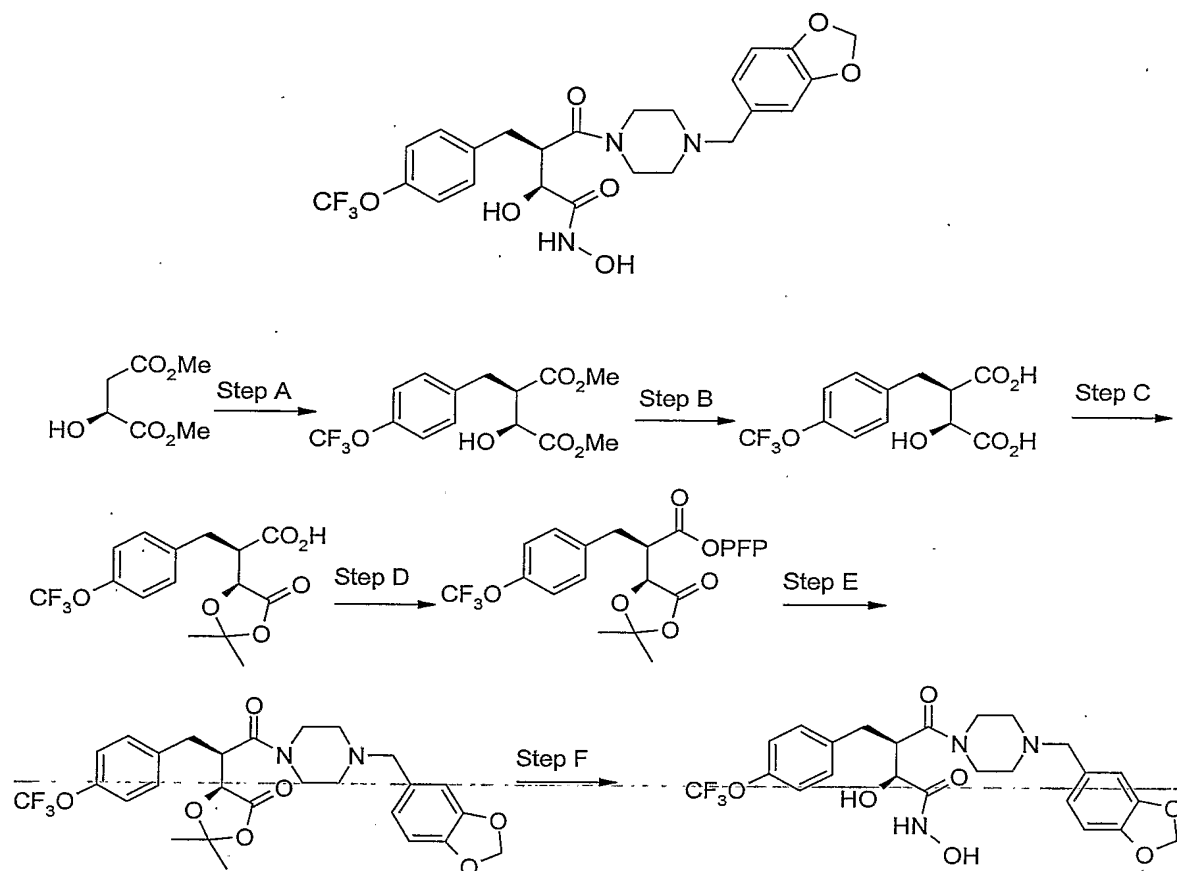
Example 28: 3R-(4-benzyl-3RS-phenyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



LRMS: +ve ion 546 (M+H).

Example 29: 4-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-2S,N-dihydroxy-4-oxo-3R-(4-trifluoromethoxy-benzyl)-butyramide.

Example 29 was prepared as outlined in Scheme 4 using procedures described below.



Reagents and conditions. A: LiHMDS, 4-OCF₃-C₆H₄-CH₂Br, THF, -78 to RT; B: NaOH, THF, H₂O; C: CuCl₂, dimethoxyacetone, acetone; D: pentafluorophenol, WSC, HOAt, CH₂Cl₂; E: RNH₂, NEt₃CH₂Cl₂; F: HONH₂ aq., iPrOH

Scheme 4

Step A: 2S-hydroxy-3R-(4-trifluoromethoxy-benzyl)-succinic acid dimethyl ester.

To a cold (-78C) solution of 2S-hydroxy-succinic acid dimethyl ester (3.0 g, 18.5 mmol) in THF (100 ml), was added a 1N solution of LiHMDS (40.7 ml, 40.7 mmol, 2.2 eq.) dropwise. The reaction mixture was stirred at -78C for 30 min and then at -30C for 1 hr. The temperature was brought down to -78C and 1-bromomethyl-4-trifluoromethoxy-benzene (3.11 ml, 19.4 mmol, 1.05 eq.) was added dropwise. The reaction mixture was allowed to warm to RT overnight and was then poured into NH₄Cl_{sat} (50 ml). THF was removed under



vacuo and the crude was taken-up in AcOEt (150 ml). The organic layer was washed with water (2 X 20 ml) and with brine (20 ml) before being dried over MgSO_4 . Solvent was removed under reduced pressure to give an oil which was purified by flash chromatography on silica gel. The expected 2S-hydroxy-3R-(4-trifluoromethoxy-benzyl)-succinic acid dimethyl ester was obtained in 24% yield (1.50 g).

$^1\text{H-NMR}$; delta (CDCl_3): 7.31 (2H, d, $J = 8.5\text{Hz}$), 7.15 (2H, d, $J = 8.4\text{Hz}$), 4.12 (1H, d, $J = 6.5\text{Hz}$), 3.75 (3H, s), 3.69 (3H, s), 3.20 (2H, m), 3.01 (1H, m).

LRMS: +ve ion 358 ($\text{M} + \text{Na}$).

Step B: 2S-hydroxy-3R-(4-trifluoromethoxy-benzyl)-succinic acid.

To a cold (0C) solution of 2S-hydroxy-3R-(4-trifluoromethoxy-benzyl)-succinic acid dimethyl ester (1.50 g, 4.45 mmol) in MeOH/water (3:1, 28 ml) was added lithium hydroxide (617 mg, 14.7 mmol, 3.3 eq.). The reaction mixture was stirred at RT for 16 hrs. The reaction was quenched by addition of HCl 1N (5 ml) and MeOH was removed under reduced pressure. The extraction was carried out with AcOEt (10 ml). The organic layer was washed with HCl 1N (10 ml), water (10 ml) and finally with brine (10 ml). The organic layer was dried over MgSO_4 . Solvent was removed under reduced pressure to give the expected 2S-hydroxy-3R-(4-trifluoromethoxy-benzyl)-succinic acid as a clear oil (1.36 g, 99% yield).

$^1\text{H-NMR}$; delta (CDCl_3): 7.32 (2H, d, $J = 8.4\text{Hz}$), 7.19 (2H, d, $J = 8.4\text{Hz}$), 4.08 (1H, d, $J = 6.5\text{Hz}$), 3.25 (2H, m), 3.05 (1H, m).

LRMS: +ve ion 331 ($\text{M} + \text{Na}$), -ve ion 307 ($\text{M} - \text{H}$).

Step C: 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-3-(4-trifluoromethoxy-phenyl)-propionic acid.

To a solution of 2S-hydroxy-3R-(4-trifluoromethoxy-benzyl)-succinic acid (1.36 g, 4.4 mmol) in acetone (10 ml) under an inert atmosphere were added dimethoxy propane (923 μl , 7.5 mmol, 1.7 eq.) and copper chloride (59 mg, 0.44 mmol, 0.1 eq.). The reaction mixture was stirred at RT for 16 hrs. The

solvent was then removed under vacuo to give 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-3-(4-trifluoromethoxy-phenyl)-propionic acid (1.08 g, 70% yield).

$^1\text{H-NMR}$; δ (CDCl_3): 7.29 (2H, d, $J = 8.7\text{Hz}$), 7.14 (2H, d, $J = 8.4\text{Hz}$), 4.30 (1H, d, $J = 6.2\text{Hz}$), 3.21 (2H, m), 3.0 (1H, m), 1.61 (3H, s), 1.52 (3H, s).

LRMS: +ve ion 371 ($\text{M} + \text{Na}$), -ve ion 347 ($\text{M} - \text{H}$).

Step D: 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-3-(4-trifluoromethoxy-phenyl)-propionic acid pentafluorophenyl ester.

To a cold (0°C) solution of 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-3-(4-trifluoromethoxy-phenyl)-propionic acid (1.08 g, 3.1 mmol) and pentafluorophenol (685 mg, 3.7 mmol, 1.2 eq.) in CH_2Cl_2 (10 ml) was added WSC (714 mg, 3.7 mmol, 1.2 eq.). The reaction mixture was allowed to warm to RT overnight. CH_2Cl_2 was removed under vacuo and the resulting crude reaction mixture was dissolved in AcOEt (80 ml). The organic layer was washed with water (20 ml), NaHCO_3 sat (10 ml) and finally with brine (10 ml). Solvent was removed under reduced pressure to give an oil which was purified by flash chromatography to furnish the expected 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-3-(4-trifluoromethoxy-phenyl)-propionic acid pentafluorophenyl ester (254 mg, 16% yield).

$^1\text{H-NMR}$; δ (CDCl_3): 7.29 (2H, bd, $J = 9.3\text{Hz}$), 7.21 (2H, bd, $J = 8.7\text{Hz}$), 4.45 (1H, d, $J = 6.2\text{Hz}$), 3.51 (1H, m), 3.3 (2H, m), 1.65 (3H, s), 1.58 (1H, s).

LRMS: +ve ion 537 ($\text{M} + \text{Na}$).

Step E: 5S-[2-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-2-oxo-1R-(4-trifluoromethoxy-benzyl)-ethyl]-2,2-dimethyl-[1,3]dioxolan-4-one

To a solution of 2-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-3R-(4-trifluoromethoxy-phenyl)-propionic acid pentafluorophenyl ester (59 mg, 0.11 mmol) in CH_2Cl_2 (10 ml) was added 1-benzo[1,3]dioxol-5-ylmethyl-piperazine (28 mg, 0.13 mmol, 1.1 eq.). The reaction mixture was stirred for 16 hrs and the solvent was removed under vacuo. The crude was taken-up in AcOEt (50 ml) and washed with water (10 ml), then with NaHCO_3 sat (10 ml) and finally with brine (10 ml). The solvent was dried over MgSO_4 and removed under

reduced pressure to give an oil which was purified by preparative reverse phase chromatography to afford the expected 5S-[2-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-2-oxo-1R-(4-trifluoromethoxy-benzyl)-ethyl]-2,2-dimethyl-[1,3]dioxolan-4-one.

¹H-NMR; delta (CD₃OD): 7.27-7.12 (4H,m), 6.78-6.62 (3H, m), 5.93 (2H, s), 4.57 (1H, d, J=6.6 Hz), 3.85-2.9 (10H, m), 2.44-2.21 (2H, m), 2.05-1.91 (1H, m), 1.69-1.42 (6H, m).

LRMS: +ve ion 551 (M+H).

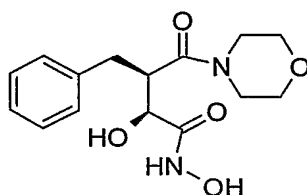
Step F: 4-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-2S,N-dihydroxy-4-oxo-3R-(4-trifluoromethoxy-benzyl)-butyramide.

To a solution of 5S-[2-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-2-oxo-1R-(4-trifluoromethoxy-benzyl)-ethyl]-2,2-dimethyl-[1,3]dioxolan-4-one (26 mg, 0.047 mmol) in *i*-PrOH (2 ml), was added an aqueous solution of hydroxylamine (50%, 16 μ l, 0.24 mmol, 5 eq.). The reaction mixture was allowed to stir at RT for 16 hrs. The solvent was removed under reduced pressure to yield an oil which was purified by preparative reverse phase chromatography to give the required 4-(4-Benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-2S,N-dihydroxy-4-oxo-3R-(4-trifluoromethoxy-benzyl)-butyramide.

¹H-NMR; delta (CD₃OD): 7.32-7.15 (4H), 6.79-6.65 (4H, m), 5.90 (2H, s), 4.13 (1H, d, J=6.8Hz), 3.75-3.35 (4H, m), 3.30 (2H, s), 3.0-2.8 (4H, m), 2.5-2.25 (2H, m), 1.90 (1H,m).

LRMS: +ve ion 526 (M+H); -ve ion 524 (M-H).

Example 30: 3R-benzyl-2S,N-dihydroxy-4-morpholin-4-yl-4-oxo-butamide.

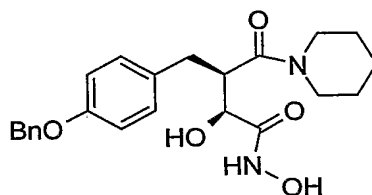


The compound of example 30 was prepared by the method of example 29 according to scheme 4, and using the benzyl bromide in step A and morpholine in step E.

$^1\text{H-NMR}$; δ (CD_3OD): 7.32-7.18 (5H, m), 4.16 (1H, d, $J=6.7\text{Hz}$), 3.59-2.67 (11H, m).

LRMS: +ve ion 331 ($\text{M}+\text{Na}$); -ve ion 307 ($\text{M}-\text{H}$).

Example 31: 3R-(4-Benzyloxy-benzyl)-2S,N-dihydroxy-4-oxo-4-piperidin-1-yl-butamide.

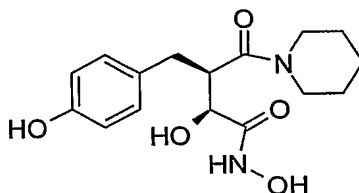


The compound of example 31 was prepared by the method of example 29 according to scheme 4, and using the 1-benzyloxy-4-bromomethyl-benzene in step A and piperidine in step E.

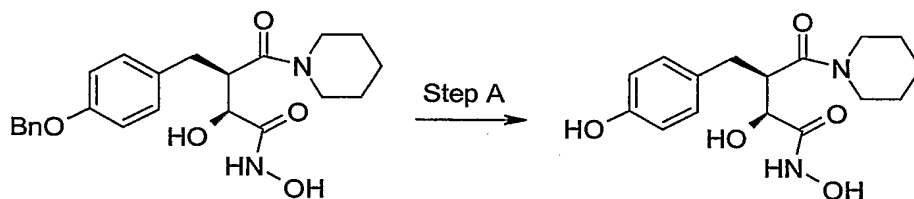
$^1\text{H-NMR}$; δ (MeOD): 7.41-7.26 (5H, m), 7.10-6.87 (4H, m), 5.05 (2H, s), 4.11 (1H, d, $J=6.1\text{Hz}$), 3.57-3.48 (2H, m), 3.24-3.09 (3H, m), 2.89-2.69 (2H, m), 1.41-1.17 (5H, m) and 0.72-0.67 (1H, m).

LRMS: +ve ion 435 ($\text{M}+\text{Na}$); -ve ion 411 ($\text{M}-\text{H}$).

Example 32: 2S,N-dihydroxy-3R-(4-hydroxy-benzyl)-4-oxo-4-piperidin-1-yl-butamide.



Example 32 was prepared as outlined in Scheme 5 using procedures described below.



Reagents and conditions: A: H_2 , Pd/C, MeOH

Scheme 5

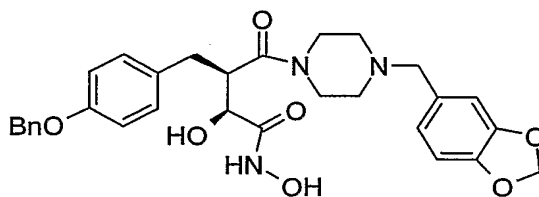
Step A: 2S,N-dihydroxy-3R-(4-hydroxy-benzyl)-4-oxo-4-piperidin-1-yl-butyramide.

To a solution of 2S,N-dihydroxy-3R-(4-hydroxy-benzyl)-4-oxo-4-piperidin-1-yl-butyramide (14 mg, $3.5 \cdot 10^{-5}$ mol) in MeOH (5 ml) under an inert atmosphere, was added 10%Pd/C (1 mg). H_2 was then bubbled through the resulting suspension for 2 hrs. Pd/C was filtered off and the solvent was removed under reduced pressure to give 2S,N-dihydroxy-3R-(4-hydroxy-benzyl)-4-oxo-4-piperidin-1-yl-butyramide (10 mg, quant.).

1H -NMR; delta (MeOD): 7.02-6.67 (4H, 2d, $J=8.4\text{Hz}$ and $J\ 8.4\text{Hz}$), 4.12 (1H, d, $J=6.0\text{Hz}$), 3.65-3.48 (2H, m), 3.27-3.12 (3H, m), 2.87-2.68 (2H, m), 1.47-1.26 (5H, m) and 0.83-0.78 (1H, m).

LRMS: +ve ion 345 (M+Na); -ve ion 321 (M-H).

Example 33: 4-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-3R-(4-benzyloxy-benzyl)-2S,N-dihydroxy-4-oxo-butyramide.



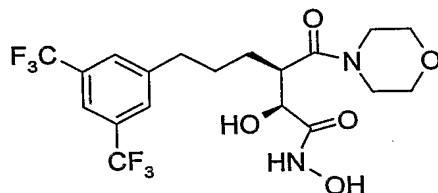
The compound of example 33 was prepared by the method of example 29 according to scheme 4, and using the 1-benzyloxy-4-bromomethyl-benzene in step A.

$^1\text{H-NMR}$; delta (DMSO): 9.80 (1H, s), 8.88 (1H, s), 7.48-7.31 (5H, m), 7.04-6.90 (4H, m), 6.82-6.77 (2H, m), 6.67-6.62 (1H, m), 5.98 (2H, s), 5.45 (1H, d), 5.09 (2H, s), 3.91 (1H, m), 3.56-3.44 (2H, m), 3.20-3.08 (6H, m), 2.78-2.60 (1H, m), 2.30-2.08 (2H, m), 1.92-1.83 (1H, m) and 1.42-1.30 (1H, m).

LRMS: +ve ion 548 (M+H); -ve ion 546 (M-H).

Example 34: 6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

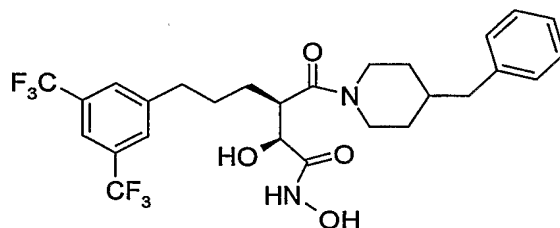
The compounds of Examples 34-38 were prepared by the method of Example 1 according to scheme 1, where ArBr in Step B was 1-bromomethyl-3,5-bis-trifluoromethyl-benzene by parallel synthesis and using the appropriate amine in Step G. The products were purified by preparative HPLC.



$^1\text{H-NMR}$; delta (CD_3OD): 7.79 (2H, s), 7.77 (1H, s), 4.07 (1H, d, $J = 6.8$ Hz), 3.19-3.76 (9H, m), 2.73-2.98 (2H, m), 1.52-1.88 (4H, m).

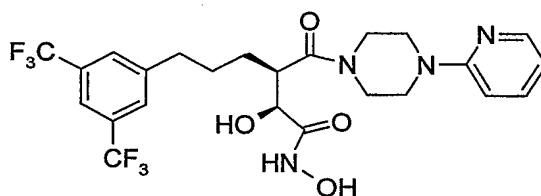
LRMS: +ve ion 473 (M+H), 495 (M+Na); -ve ion 471 (M-H).

Example 35: 3R-(4-benzyl-piperidine-1-carbonyl)-6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



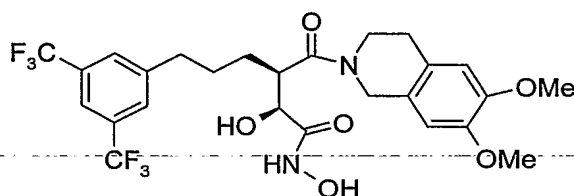
LRMS: +ve ion 561 (M+H), 583 (M+Na); -ve ion 559 (M-H).

Example 36: 6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.



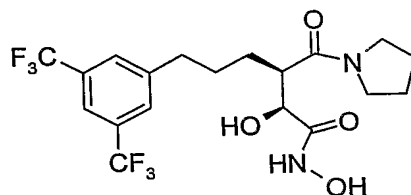
LRMS: +ve ion 549 (M+H); -ve ion 547 (M-H).

Example 37: 6-(3,5-bis-trifluoromethyl-phenyl)-3R-(6,7-dimethoxy-3,4-dihydro-1H-isoquinoline-2-carbonyl)-2S-hydroxy-hexanoic acid hydroxyamide.



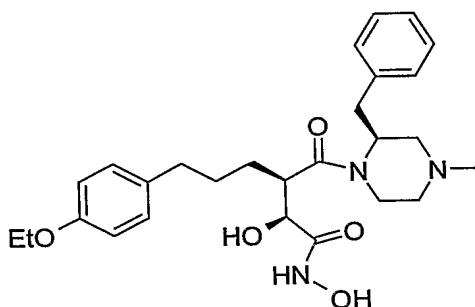
LRMS: +ve ion 601 (M+Na); -ve ion 577 (M-H).

Example 38: 6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(pyrrolidine-1-carbonyl)-hexanoic acid hydroxyamide.



$^1\text{H-NMR}$; delta (CD_3OD): 7.84 (2H, s), 7.79 (1H, s), 4.06 (1H, d, $J = 6.6$ Hz), 3.71-3.81 (1H, m), 3.10-3.54 (5H, m), 2.75-2.83 (1H, m), 1.51-1.98 (7H, m).
LRMS: +ve ion 479 ($\text{M}+\text{Na}$); -ve ion 455 ($\text{M}-\text{H}$).

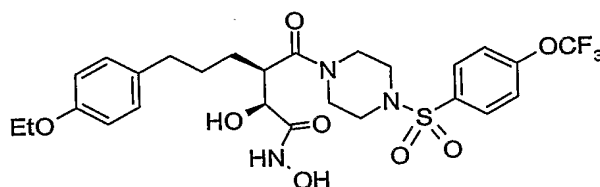
Example 39: 3R-(2S-benzyl-4-methyl-piperazine-1-carbonyl)-6-(4-ethoxyphenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

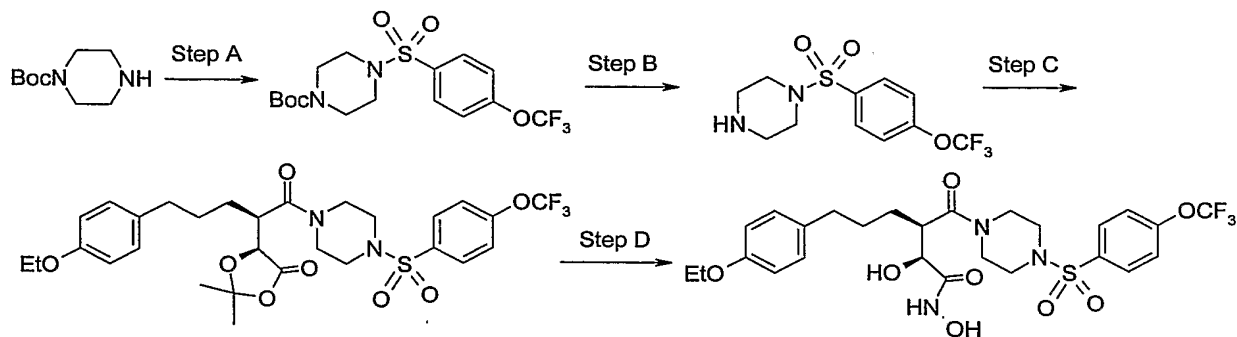


The compound of example 39 was prepared by the method of example 1 according to scheme 1, and using the appropriate piperazine (synthesised according to the procedure described in J. Heterocyclic Chem, **28**, 1219-1224, 1991; in Step A. The product was purified by preparative HPLC.

LRMS: +ve ion 484 ($\text{M}+\text{H}$).

Example 40: 6-(4-ethoxyphenyl)-2S-hydroxy-3R-[4-(4-trifluoromethoxybenzenesulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.





Reagents and conditions: A: RSO_2Cl 1.2 eq., NEt_3 1.4 eq., CH_2Cl_2 ; B: $\text{TFA}/\text{CH}_2\text{Cl}_2$; C: LHS, EDAC, HOAt, CH_2Cl_2 ; D: H_2NOH aq., iPrOH

Scheme 6

Example 40 was prepared as outlined in Scheme 6 using procedures described below.

Step A: 4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carboxylic acid tert-butyl ester.

To a solution of piperazine-1-carboxylic acid tert-butyl ester (521 mg, 2.8 mmol) in DCM (6 ml) was added NEt_3 (547 μl , 3.92 mmol, 1.4 eq.) and 4-trifluoromethoxy-benzenesulfonyl chloride (880 mg, 3.36 mmol, 1.2 eq.). The reaction mixture was stirred at room temperature for 12 hrs. The excess of sulfonyl chloride was quenched by addition of trisamine resin (150 mg), and subsequent stirring for 2 hrs. The resin was then filtered off, and NaHCO_3 sat (5 ml) was added. Filtration of the resulting mixture through hydrophobic cartridge and subsequent removal of the solvent under reduced pressure afforded the desired 4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carboxylic acid tert-butyl ester in quantitative yield.

Step B: 1-(4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine.

To a cold (0 C) solution of 4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carboxylic acid tert-butyl ester (1.15 g, 2.8 mmol) in DCM (3 ml) was added TFA (2 ml). The resulting solution was stirred for 1 hr. Solvent was then removed under reduced pressure. The crude solid was taken-up in AcOEt (15 ml) and washed with NaHCO_3 sat (2 X 5 ml), water (5 ml) and finally with brine (5 ml). The organic layer was dried over MgSO_4 and the solvent was removed

under vacuo to yield the desired 1-(4-trifluoromethoxy-benzenesulfonyl)-piperazine as a white solid (753 mg, 87% yield).

$^1\text{H-NMR}$; δ (CDCl_3): 7.80 (2H, m), 7.38 (2H, m), 3.01 (8H, m).

LRMS: +ve ion 311 (M+H).

Step C: 5-{4-(4-ethoxy-phenyl)-1-[4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-butyl}-2,2-dimethyl-[1,3]dioxolan-4-one.

To a cold (0 C) solution of 2R-(2,2-dimethyl-5-oxo-[1,3]dioxolan-4S-yl)-5-(4-ethoxy-phenyl)-pentanoic acid (91 mg, 0.27 mmol), 1-(4-trifluoromethoxy-benzenesulfonyl)-piperazine (100 mg, 0.325 mmol, 1.2 eq.) in DCM (6 ml), was added WSCDI (62 mg, 0.325 mmol, 1.2 eq.) and HOAt (cat). The reaction mixture was stirred for 16 hrs allowing the temperature to raise to 20 C. DCM was then removed under reduced pressure, and the resulting crude reaction mixture was taken-up in AcOEt (30 ml). The organic layer was washed with water (2 X 10 ml) and with brine (10 ml). The organic layer was dried over MgSO_4 and the solvent was removed under vacuo to yield the desired 5-{4-(4-ethoxy-phenyl)-1-[4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-butyl}-2,2-dimethyl-[1,3]dioxolan-4-one (170 mg, quant.).

LRMS: +ve ion 629 (M+H), 651 (M+Na).

Step D: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

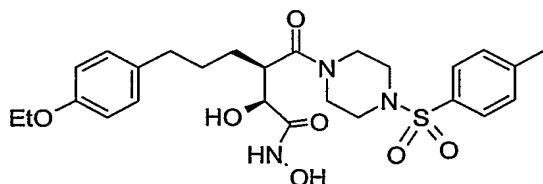
To a solution of 5-{4-(4-ethoxy-phenyl)-1-[4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-butyl}-2,2-dimethyl-[1,3]dioxolan-4-one (170 mg, 0.27 mmol) in iPrOH (5 ml) was added $\text{H}_2\text{NHOH}_{\text{aq}}$ (90 μl , 1.35 mmol, 5 eq.). The reaction mixture was stirred for 12 hrs and the solvent was removed under vacuo. The crude reaction mixture was purified through preparative HPLC to give the expected 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-

(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

$^1\text{H-NMR}$; delta (CDCl_3): 7.79 (2H, m), 7.38 (2H, m), 7.01 (2H, m), 6.80 (2H, m), 4.06 (2H, q), 3.90 (1H, m), 3.64 (1H, m), 3.43-3.20 (3H, m), 3.19-3.00 (2H, t), 2.91-2.70 (2H, dt), 2.5 (2H, t), 1.8-1.5 (2H, m), 1.41 (3H, t).

LRMS: +ve ion 604 (M+H), 651 (M+Na), -ve ion (602 (M-H).

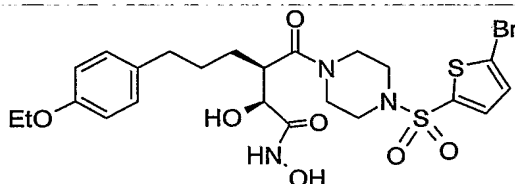
Example 41: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(toluene-4-sulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



The compound of example 41-46 were prepared by the method of example 40 according to scheme 6, and using the appropriate sulfonyl chloride in step A.

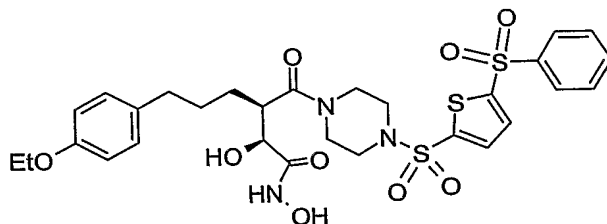
LRMS: +ve ion 534 (M+H), -ve ion 532 (M-H).

Example 42: 3R-[4-(5-bromo-thiophene-2-sulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



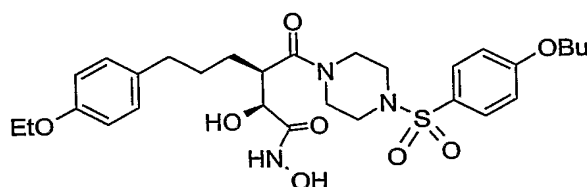
LRMS: +ve ion 605 (M+H), -ve ion 603 (M-H).

Example 43: 3R-[4-(5-benzenesulfonyl-thiophene-2-sulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



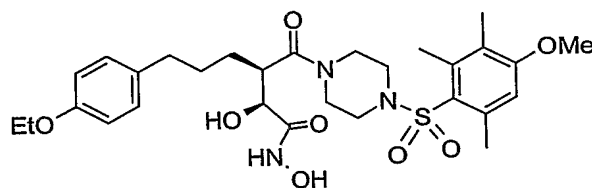
LRMS: +ve ion 666 (M+H), -ve ion 664 (M-H).

Example 44: 3R-[4-(4-butoxy-benzenesulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



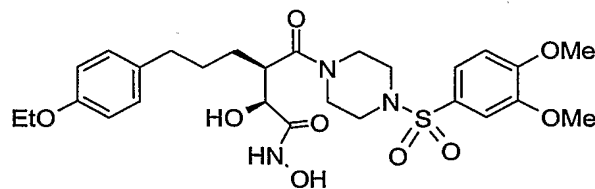
LRMS: +ve ion 592 (M+H), -ve ion 590 (M-H).

Example 45: 6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-methoxy-2,3,6-trimethyl-benzenesulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



LRMS: +ve ion 592 (M+H), -ve ion 590 (M-H).

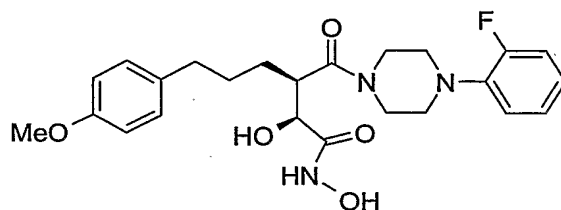
Example 46: 3R-[4-(3,4-dimethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.



LRMS: +ve ion 580 (M+H), -ve ion 578 (M-H).

The compounds of Examples 47-50 were prepared by the method of Example 1 by parallel synthesis, using the appropriate aryl bromide in Step B, and the appropriate amine in Step G. The products were purified by preparative HPLC

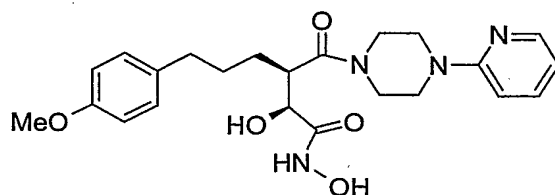
Example 47: 6-(4-methoxy-phenyl)-2S-hydroxy-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.



$^1\text{H-NMR}$; delta (MeOD): 7.20-7.00 (6H, m), 6.85 (2H, m), 4.15-4.10 (1H, m), 3.95-3.70 (6H, m), 3.45-3.35 (4H, m), 3.20-2.95 (3H, m), 2.65-2.55 (2H, m), 1.80-1.55 (3H, m).

LRMS: +ve ion 460 (M+H); -ve ion 458 (M-H).

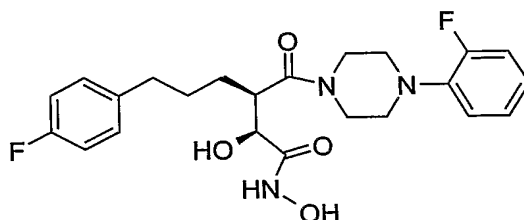
Example 48: 6-(4-methoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.



$^1\text{H-NMR}$; delta (MeOD): 8.15-8.10 (1H, m), 7.65-7.55 (1H, m), 7.15-7.05 (2H, m), 6.90-6.80 (3H, m), 6.80-6.70 (1H, m), 4.20-4.10 (1H, m), 3.95-3.35 (13H, m), 2.65-2.55 (2H, m), 1.80-1.55 (3H, m).

LRMS: +ve ion 443 (M+H).

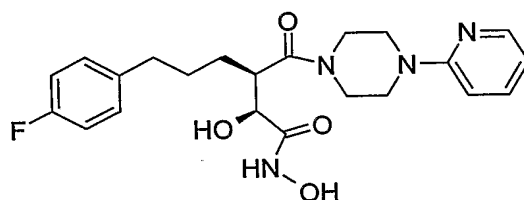
Example 49: 6-(4-fluoro-phenyl)-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-2S-hydroxy-hexanoic acid hydroxyamide.



$^1\text{H-NMR}$; delta (MeOD): 7.2-6.9 (8H, m), 4.05 (1H, d), 3.9-3.6 (4H, m), 3.35 (2H, m), 3.1-2.9 (3H, m), 2.55 (2H, t), 1.8-1.5 (3H, m), 1.15 (1H, t).

LRMS: +ve ion 448 (M+H).

Example 50: 6-(4-fluoro-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.



$^1\text{H-NMR}$; delta (MeOD): 8.1 (1H, d), 7.55 (1H, t), 7.15 (2H, m), 6.9 (2H, m), 6.8 (1H, d), 6.7 (1H, m), 4.05 (1H, d), 3.85 (2H, m), 3.7-3.3 (6H, m), 3.05 (1H, t), 2.55 (2H, t), 1.8 (4H, m), 1.15 (1H, t).

LRMS: +ve ion 431 (M+H).

Biological Results

A. Enzyme Inhibition Assays

Compounds of the invention were tested to assess their activities as inhibitors of MMP9, MMP12 and MMP-1.

MMP9 Assay Protocol

Compounds of the invention were tested for inhibitory activity against 92kDa gelatinase (MMP9) in an assay using a coumarin-labelled peptide substrate, (7-methoxycoumarin-4-yl)acetyl-Pro-Leu-Gly-Leu-(3-[2,4-dinitrophenyl]-L-2,3-diaminopropionyl)-Ala-Arg-NH₂ (McaPLGLDpaAR) (Knight et al, FEBS Lett. 1992; 263-266).

Stock solutions were made up as follows:

Assay Buffer: 100mM Tris-HCl pH 7.6 containing 100mM NaCl, 10mM CaCl₂, and 0.05% Brij 35

Substrate: 0.4mM McaPLGLDpaAR (from Bachem) (0.437mg/ml) stock solution in 100%

DMSO (stored at -20°C). Dilute to 8μM in assay buffer.

Enzyme: Recombinant human 92kDa gelatinase (MMP-9; APMA (4-aminophenylmercuric acetate) -activated if necessary) appropriately diluted in assay buffer.

Test Compounds were prepared initially as 10mM compound solution in 100% DMSO, diluted to 1mM in 100% DMSO, then serially diluted 3-fold in 100% DMSO across columns 1-10 of a 96-well microtitre plate Assay concentration range, 100μM (column 1) to 5.1nM (column 10)

The assay was performed in a total volume of 100μl per well in 96-well microtitre plates. Activated enzyme (20μl) was added to the wells followed by 20μl of assay buffer. Appropriate concentrations of test compounds dissolved in 10μl of DMSO were then added followed by 50μl of McaPLGLDpaAR (8μM, prepared by dilution of DMSO stock in assay buffer). For each assay ten

concentrations of test compound were examined in duplicate. Control wells lack either enzyme or test compound. The reactions were incubated at 37°C for 2 hours. The fluorescence at 405nm was measured immediately with an SLT Fluostar fluorometer (SLT Labinstruments GmbH, Grödig, Austria) using 320nm excitation, without stopping the reaction.

The effect of the test compound was determined from the dose response curve generated by the 10 duplicate concentrations of inhibitor. The IC₅₀ (the concentration of compound required to give a 50% decrease in enzyme activity) was obtained by fitting data to the equation, $Y = a + ((b - a) / (1 + (c/X)^d))$. (Y = inhibition achieved for a particular dose; X = the dose in nM; a = minimum y or zero % inhibition; b = maximum y or 100% inhibition; c = is the IC₅₀; d = is the slope). The result was rounded to one significant figure.

MMP12 Assay protocol

Compounds of the invention were tested for inhibitory activity against metalloelastase (MMP12) in an assay using a coumarin-labelled peptide substrate, (7-methoxycoumarin-4-yl)acetyl-Pro-Leu-Gly-Leu-(3-[2,4-dinitrophenyl]-L-2,3-diaminopropionyl)-Ala-Arg-NH₂ (McaPLGLDpaAR) (Knight et al, FEBS Lett. 1992; 263-266). The protocol for this assay was as described for the MMP9 assay above.

MMP1 Assay protocol

Compounds of the invention were tested for inhibitory activity against collagenase (MMP1) in an assay using a coumarin-labelled peptide substrate, (7-methoxycoumarin-4-yl)acetyl-Pro-Leu-Gly-Leu-(3-[2,4-dinitrophenyl]-L-2,3-diaminopropionyl)-Ala-Arg-NH₂ (McaPLGLDpaAR) (Knight et al, FEBS Lett. 1992; 263-266). The protocol for this assay was as described for the MMP9 assay above.

Results:

Key to assay data

Range	A	< 100nM
	B	100 – 1000nM

C	1000 – 10,000nM
D	>10,000nM

For instance the inhibitory activities of some of the compounds of the examples against MMP-12, MMP-9, MMP-2 and MMP-1 are set out in the following table:

Example Number	MMP-9 (nM)	MMP-2 (nM)	MMP-1 (nM)	MMP-12 (nM)
4	A	A	D	A
5	A	A	D	A
6	B	B	D	A
12	A	A	D	A
13	A	A	D	A
14	A	A	D	A
20	A	A	D	A

B. Biological Assay: IL2-induced peritoneal recruitment of lymphocytes in mice

Protocol

C3H Mice (Elevage Janvier) (8 week old, n=6) received IL2 (SPRI, 20 µg/kg, 40 ml/kg, ip) 15 min after administration of the test molecules. Twenty-four hours later, the animals were sacrificed and the lavage of the peritoneal cavity was conducted using 3 x 5 ml PBS-1mM EDTA (+4°C). After centrifugation (10 min at 3000 rpm), the pellet was resuspended in 1 ml PBS. The peritoneal cells were counted using a Beckman/Coulter counter. IL2 was solubilized in saline. The test compounds of the invention were solubilized or suspended in 0.5% carboxymethylcellulose (CMC)/0.25% Tween-20 and orally administered. Dexamethasone (0.1 mg/kg, po) was used as reference compound.

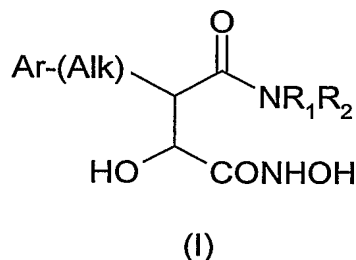
For instance, upon using the compounds of specified examples, the inhibition of IL-2 induced peritoneal recruitment was as set out in the following table:

IL2-Induced peritoneal Recruitment

	Doses (mg/kg)	Route	% Inhibition
Example 4	3	po	34%
Example 5	3	po	50%
Example 6	3	po	33%
Example 13	3	po	47%
Example 14	3	po	30%
Example 20	3	po	57%
Dexamethazone	1	sc	76%

Claims:

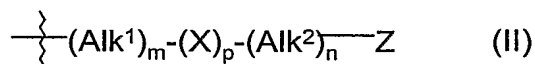
1. A compound of formula (I), or an enantiomer or diastereoisomer thereof, or a salt, hydrate or solvate thereof:

wherein

Ar represents an optionally substituted aryl, heteroaryl, C₃-C₈ cycloalkyl or heterocycloalkyl group;

Alk represents a divalent C₁-C₅ alkylene or C₂-C₅ alkenylene radical; and

R₁ and R₂ taken together with the nitrogen atom to which they are attached form a first heterocycloalkyl ring which is optionally fused to a second C₃-C₈ cycloalkyl or heterocycloalkyl ring, the said first and second rings being optionally substituted by at least one group of formula (II):

wherein

m, p and n are independently 0 or 1;

Z represents, hydrogen, or an optionally substituted carbocyclic or heterocyclic ring of from 5 to 7 ring atoms which is optionally fused to another optionally substituted carbocyclic or heterocyclic ring of from 5 to 7 ring atoms;

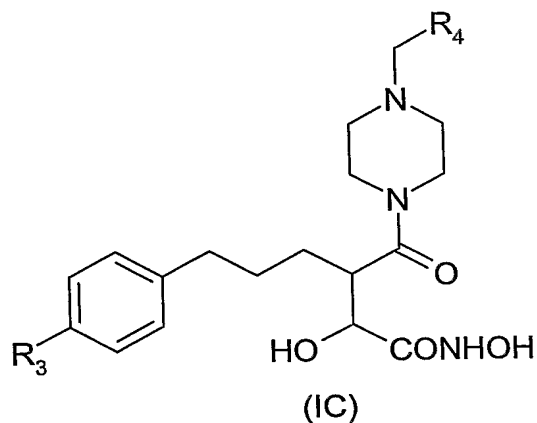
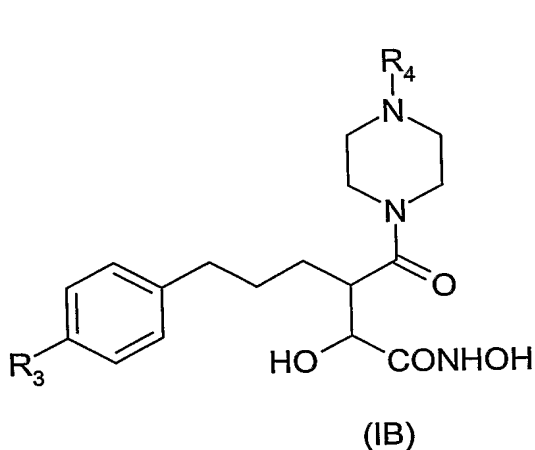
Alk¹ and Alk² independently represent optionally substituted divalent C₁-C₃ alkylene radicals;

X represents -O-, -S-, -S(O)-, -S(O₂)-, -C(=O)-, -NH-, -NR₃-,

-S(O₂)NH-, -S(O₂)NR₃-, -NHS(O₂)-, or -NR₃S(O₂)-, where R₃ is C₁-C₃ alkyl.

2. A compound as claimed in claim 1 wherein Ar is a 5- or 6- membered monocyclic aryl or heteroaryl ring, which is optionally substituted by at least one substituent selected from (C₁-C₃)alkyl, (C₁-C₃)alkoxy, hydroxy, hydroxy(C₁-C₃)alkyl, mercapto, mercapto(C₁-C₃)alkyl, (C₁-C₃)alkylthio, halo, trifluoromethyl, trifluoromethoxy, nitro, nitrile (-CN), -COOH, -COOR^A, -COR^A, -SO₂R^A, -CONH₂, -SO₂NH₂, -CONHR^A, -SO₂NHR^A, -CONR^AR^B, -SO₂NR^AR^B, -NH₂, -NHR^A, -NR^AR^B, -OCONH₂, -OCONHR^A, -OCONR^AR^B, -NHCOR^A, -NHCOOR^A, -NR^BCOOR^A, -NHSO₂OR^A, -NR^BSO₂OR^A, -NHCONH₂, -NR^ACONH₂, -NHCONHR^B, -NR^ACONHR^B, -NHCONR^AR^B, or -NR^ACONR^AR^B wherein R^A and R^B are independently C₁-C₃ alkyl, phenyl or a 5- or 6- membered monocyclic aryl or heteroaryl ring.
3. A compound as claimed in claim 2 wherein an optional substituent is in the 4- position in the case of a 6-membered ring, or in the 2- and/or 3- position in the case of a 5-membered ring.
4. A compound as claimed in any of the preceding claims wherein Ar is optionally substituted phenyl, 2-, 3-, or 4-pyridyl, 2-, or 3-thienyl, or 2-, or 3-furanyl.
5. A compound as claimed in any of the preceding claims wherein optional substituents in Ar are selected from methoxy, ethoxy, trifluoromethoxy, methyl, ethyl, trifluoromethyl, hydroxyl, mercapto, fluoro, chloro, and bromo.
6. A compound as claimed in claim 2 wherein Ar is 4-(C₁C₃alkoxy)phenyl.
7. A compound as claimed in claim 2 wherein Ar is 4-ethoxyphenyl
8. A compound as claimed in any of the preceding claims wherein Alk is -CH₂-, -CH₂CH₂-, -CH₂CH(CH₃)-, -CH₂CH₂CH₂-, -CH₂CH₂CH₂CH₂-, -CH=CH-, -CH₂CH=CH-, -CH₂CH=CHCH₂-, or -CH=CHCH=CH-.

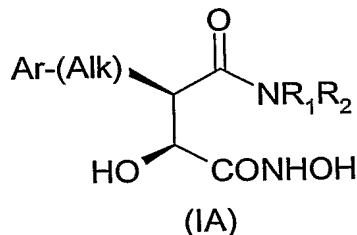
9. A compound as claimed in any of the preceding claims wherein $-NR_1R_2$ forms a pyrrolidinyl, morpholyl, or thiomorpholyl ring
10. A compound as claimed in any of claims 1 to 8 wherein $-NR_1R_2$ forms a piperidinyl, or piperazinyl ring
11. A compound as claimed in any of the preceding claims wherein in the group (II), when present, p is 0, Z is hydrogen and at least one of n and m is 1.
12. A compound as claimed in any of claims 1 to 10 wherein in the group (II), when present, m, n and p are all 0 and Z is a carbocyclic or heterocyclic ring directly linked to a ring carbon or ring nitrogen of the $-NR_1R_2$ group.
13. A compound as claimed in any of claims 1 to 10 wherein in the group (II), when present, p is 0, at least one of m and n is 1, and Z is a carbocyclic or heterocyclic ring linked to a ring carbon or ring nitrogen of the $-NR_1R_2$ group via a C_1-C_6 alkylene linker between Z and the $-NR_1R_2$ ring.
14. A compound as claimed in any of claims 1 to 10 wherein in the group (II), when present, p is 1.
15. A compound as claimed in claim 1 or claim 2 of formula (IB) or (IC) or an enantiomer or diastereoisomer thereof, or a salt, hydrate or solvate thereof:



wherein R_3 is trifluoromethyl, trifluoromethoxy C_1 - C_3 alkoxy, hydroxy, or halo; R_4 is (i) $-SO_2R_5$ or $-COR_5$ wherein R_5 is C_1 - C_6 alkyl or phenyl or monocyclic heteroaryl having 5 or 6 ring atoms, optionally substituted by (C_1 - C_3)alkyl, (C_1 - C_3)alkoxy, hydroxy, hydroxy(C_1 - C_3)alkyl, mercapto, mercapto(C_1 - C_3)alkyl, (C_1 - C_3)alkylthio, halo, trifluoromethyl, trifluoromethoxy or (ii) phenyl or monocyclic heteroaryl having 5 or 6 ring atoms; optionally substituted by (C_1 - C_3)alkyl, (C_1 - C_3)alkoxy, hydroxy, hydroxy(C_1 - C_3)alkyl, mercapto, mercapto(C_1 - C_3)alkyl, (C_1 - C_3)alkylthio, halo, trifluoromethyl, trifluoromethoxy.

16. A compound as claimed in claim 15 wherein a heteroaryl ring forming part of R_4 is pyridyl, pyrimidinyl, triazinyl, thienyl, or furanyl.

17. A compound as claimed in any of the preceding claims having the stereochemical configuration shown in formula (IA):



18. A compound as claimed in claim 1 or claim 2, which is selected from the group consisting of:

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(pyrrolidine-1-carbonyl)-hexanoic acid hydroxyamide.

3R-(6,7-dimethoxy-3,4-dihydro-1H-isoquinoline-2-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(3-methoxy-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-methoxy-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-4-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(2RS-methyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(2,6-RS-dimethyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(thiomorpholine-4-carbonyl)-hexanoic acid hydroxyamide.

3R-(4-benzyl-piperidine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-(4-benzo[1,3]dioxol-5-ylmethyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-4-ylmethyl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-benzylpiperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(4-pyrimidin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-trifluoromethyl-pyrimidin-2-yl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-chloro-pyrimidin-2-yl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

3R-[4-(4,6-dimethoxy-[1,3,5]triazin-2-yl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(3-trifluoromethyl-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

3R-[4-(acetyl-methyl-amino)-piperidine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(methyl-propyl-amino)-piperidine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-benzyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-isobutyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-(3S-phenyl-morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

3R-(4-benzyl-3RS-methyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-(3S,4-dibenzyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-(4-benzyl-3RS-phenyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

4-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-2S,N-dihydroxy-4-oxo-3R-(4-trifluoromethoxy-benzyl)-butyramide.

3R-benzyl-2S,N-dihydroxy-4-morpholin-4-yl-4-oxo-butylamide.

3R-(4-Benzyloxy-benzyl)-2S,N-dihydroxy-4-oxo-4-piperidin-1-yl-butylamide.

2S,N-dihydroxy-3R-(4-hydroxy-benzyl)-4-oxo-4-piperidin-1-yl-butylamide.

4-(4-benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-3R-(4-benzyloxy-benzyl)-2S,N-dihydroxy-4-oxo-butylamide.

6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(morpholine-4-carbonyl)-hexanoic acid hydroxyamide.

3R-(4-benzyl-piperidine-1-carbonyl)-6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(3,5-bis-trifluoromethyl-phenyl)-3R-(6,7-dimethoxy-3,4-dihydro-1H-isoquinoline-2-carbonyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(3,5-bis-trifluoromethyl-phenyl)-2S-hydroxy-3R-(pyrrolidine-1-carbonyl)-hexanoic acid hydroxyamide

3R-(2S-benzyl-4-methyl-piperazine-1-carbonyl)-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-trifluoromethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(toluene-4-sulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

3R-[4-(5-bromo-thiophene-2-sulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-[4-(5-benzenesulfonyl-thiophene-2-sulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

3R-[4-(4-butoxy-benzenesulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-ethoxy-phenyl)-2S-hydroxy-3R-[4-(4-methoxy-2,3,6-trimethyl-benzenesulfonyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

3R-[4-(3,4-dimethoxy-benzenesulfonyl)-piperazine-1-carbonyl]-6-(4-ethoxy-phenyl)-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-methoxy-phenyl)-2S-hydroxy-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-hexanoic acid hydroxyamide.

6-(4-methoxy-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

6-(4-fluoro-phenyl)-3R-[4-(2-fluoro-phenyl)-piperazine-1-carbonyl]-2S-hydroxy-hexanoic acid hydroxyamide.

6-(4-fluoro-phenyl)-2S-hydroxy-3R-(4-pyridin-2-yl-piperazine-1-carbonyl)-hexanoic acid hydroxyamide.

19. A pharmaceutical composition comprising a compound as claimed in any of the preceding claims, together with a pharmaceutically acceptable carrier.

20. A compound as claimed in any of claims 1 to 18 for use as a medicament.

21. A method of treatment or prophylaxis of diseases or conditions responsive to inhibition of MMP-12 and/or MMP-9 in mammals, which method comprises administering to the mammal an effective amount of a compound as claimed in any of claims 1 to 18

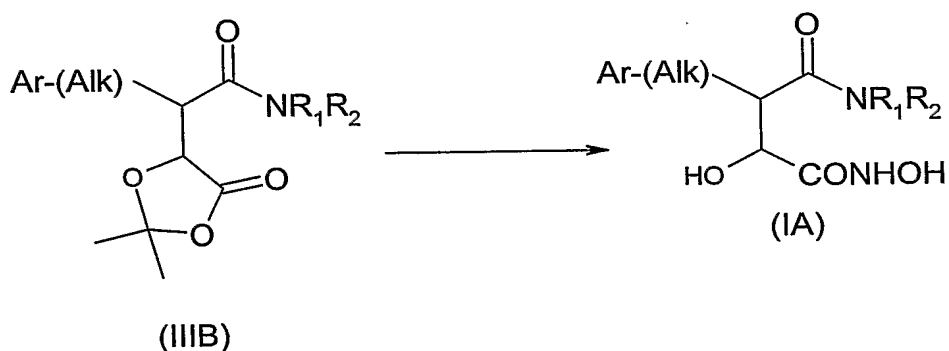
22. The use of a compound as claimed in any of claims 1 to 18 in the preparation of an agent for the treatment or prophylaxis of diseases or conditions responsive to inhibition of MMP-12 and/or MMP-9.

23. A method as claimed in claim 21 or a use as claimed in claim 22 wherein the disease or condition is bone resorption, tumour growth or invasion by secondary metastases; rheumatoid arthritis, septic arthritis, osteoarthritis, periodontitis, gingivitis, corneal ulceration, cardiac hypertrophy, acute respiratory distress syndrome, neuroinflammatory disorders, e.g. multiple sclerosis; restenosis; emphysema; fibrotic disease e.g. fibrosis post

radiotherapy, kerotid scarring, liver fibrosis and cystic fibrosis; chronic obstructive pulmonary disease; bronchitis; asthma; autoimmune disease; transplant rejection (e.g. graft versus host disease); cystic fibrosis; psoriasis; psoriatic arthritis; degenerative cartilage loss; inflammatory gastric conditions, e.g. Crohn's disease, inflammatory bowel disease, and ulcerative colitis; atopic dermatitis, epidermolysis bullosa; epidermic ulceration; a neuropathy or nephropathy e.g. interstitial nephritis, glomerulonephritis or renal failure; ocular inflammation; liver cirrhosis, Sjogren's syndrome; or an inflammatory condition of the nervous system.

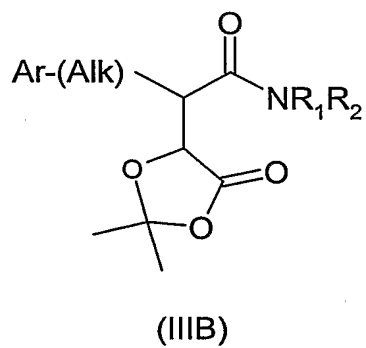
24. A method as claimed in claim 21 or a use as claimed in claim 22 wherein the disease or condition is fibrotic disease, multiple sclerosis, emphysema, bronchitis or asthma.

25. A method of preparing metalloproteinase inhibitors of formula (IA) according to any of claims 1 to 18, comprising the de-protection and/or transformation step of:



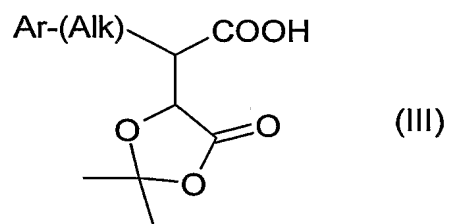
wherein Ar, Alk, R₁ and R₂ are as defined in any of claims 1 to 16.

26. A compound of formula (IIIB)



wherein Ar, Alk, R₁ and R₂ are as defined in any of claims 1 to 16.

27. A process for the preparation of a compound as claimed in claim 26 comprising comprising the step of reacting a compound of formula (III)



with a cyclic amine HNR₁R₂., wherein Ar, Alk, R₁ and R₂ are as defined in any of claims 1 to 16.

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